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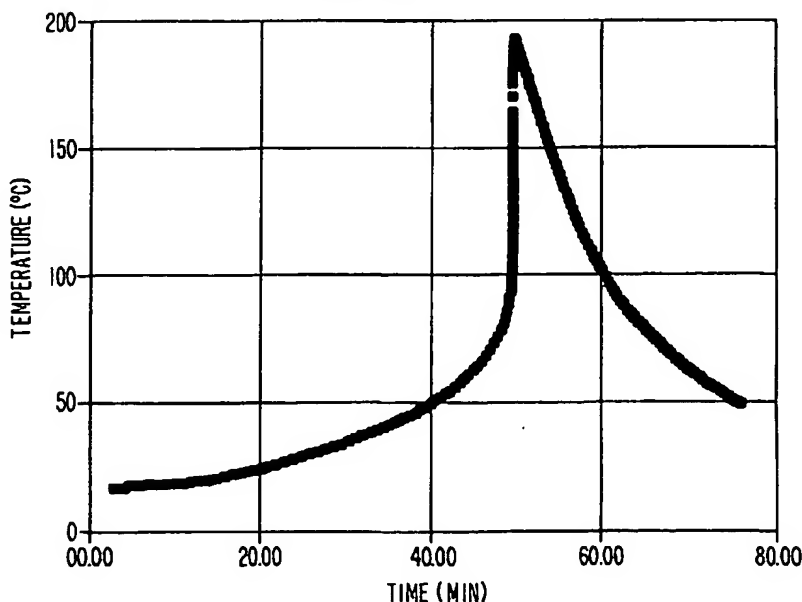
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[Continued on next page]

(54) Title: PROCESS FOR THE PREPARATION OF ANILINE-DERIVED THYROID RECEPTOR LIGANDS

REDUCED DATA



(57) Abstract: Provided are processes for the synthesis of aniline derivatives, specifically certain aniline derivatives which have activity as thyroid receptor ligands.

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PROCESS FOR THE PREPARATION OF
ANILINE-DERIVED THYROID RECEPTOR LIGANDS

5

BACKGROUND OF THE INVENTION

The present invention relates to processes for the synthesis of organic molecules, specifically molecules that have activity as thyroid receptor ligands.

10

While the essential role of thyroid hormones in regulating metabolism in humans is well recognized, the discovery and development of new specific drugs for improving the treatment of hyperthyroidism and
15 hypothyroidism has been slow. This has also limited the development of thyroid agonists and antagonists for treatment of other important clinical indications, such as hypercholesterolemia, obesity and cardiac arrhythmias.

20

Thyroid hormones affect the metabolism of virtually every cell of the body. At normal levels, these hormones maintain body weight, the metabolic rate, body temperature, and mood, and influence serum-low density lipoprotein (LDL) levels. Thus, in hypothyroidism there
25 is weight gain, high levels of LDL cholesterol, and depression. In excess with hyperthyroidism, these hormones lead to weight loss, hypermetabolism, lowering of serum LDL levels, cardiac arrhythmias, heart failure, muscle weakness, bone loss in postmenopausal women, and
30 anxiety.

Thyroid hormones are currently used primarily as replacement therapy for patients with hypothyroidism.

Therapy with L-thyroxine returns metabolic functions to normal and can easily be monitored with routine serum measurements of levels of thyroid-stimulating hormone (TSH), thyroxine (3,5,3',5'-tetraiodo-L-thyronine, or T₄) and triiodothyronine (3,5,3'-triiodo-L-thyronine, or T₃). However, replacement therapy, particularly in older individuals, is limited by certain of the deleterious effects of thyroid hormones.

In addition, some effects of thyroid hormones may be therapeutically useful in non-thyroid disorders if adverse effects can be minimized or eliminated. These potentially useful influences include weight reduction, lowering of serum LDL levels, amelioration of depression and stimulation of bone formation. Prior attempts to utilize thyroid hormones pharmacologically to treat these disorders have been limited by manifestations of hyperthyroidism, and in particular by cardiovascular toxicity.

Development of specific and selective thyroid hormone receptor agonists could lead to specific therapies for these common disorders while avoiding the cardiovascular and other toxicities of native thyroid hormones. Tissue-selective thyroid hormone agonists may be obtained by selective tissue uptake or extrusion, topical or local delivery, targeting to cells through other ligands attached to the agonist and targeting receptor subtypes. Thyroid hormone receptor agonists that interact selectively with the β -form of the thyroid hormone receptor offer an especially attractive method for avoiding cardio-toxicity.

Thyroid hormone receptors (TRs) are, like other nuclear receptors, single polypeptide chains. The various receptor forms appear to be products of two different genes α and β . Further isoform differences are due to the fact that differential RNA processing results in at least two isoforms from each gene. The $TR\alpha_1$, $TR\beta_1$ and $TR\beta_2$ isoforms bind thyroid hormone and act as ligand-regulated transcription factors. In adults, the $TR\beta_1$ isoform is the most prevalent form in most tissues, especially in the liver and muscle. The $TR\alpha_1$ isoform is prevalent in the pituitary and other parts of the central nervous system, does not bind thyroid hormones, and acts in many contexts as a transcriptional repressor. The $TR\alpha_1$ isoform is also widely distributed, although its levels are generally lower than those of the $TR\beta_1$ isoform. This isoform may be especially important for development. Whereas many mutations in the $TR\beta$ gene have been found and lead to the syndrome of generalized resistance to thyroid hormone, mutations leading to impaired $TR\alpha$ function have not been found.

A growing body of data suggest that many or most effects of thyroid hormones on the heart, and in particular on the heart rate and rhythm, are mediated through the α -form of the $TR\alpha_1$ isoform, whereas most actions of the hormone such as on the liver, muscle and other tissues are mediated more through the β -forms of the receptor. Thus, a $TR\beta$ -selective agonist might not elicit the cardiac rhythm and rate influences of the hormones but would elicit many other actions of the hormones. It is believed that the α -form of the receptor

is the major drive to heart rate for the following reasons:

- 5 1. tachycardia is very common in the syndrome of generalized resistance to thyroid hormone in which there are defective TR β -forms, and high circulating levels of T₄ and T₃;
- 10 2. there was a tachycardia in the only described patient with a double deletion of the TR β gene (Takeda et al., J. Clin. Endocrinol. & Metab. 1992, Vol. 74, p. 49);
- 15 3. a double knockout TR α gene (but not β -gene) in the mouse has a slower pulse than control mice; and,
4. western blot analysis of human myocardial TRs show presence of the TR α ₁, TR α ₂ and TR β ₂ proteins, but not TR β ₁.

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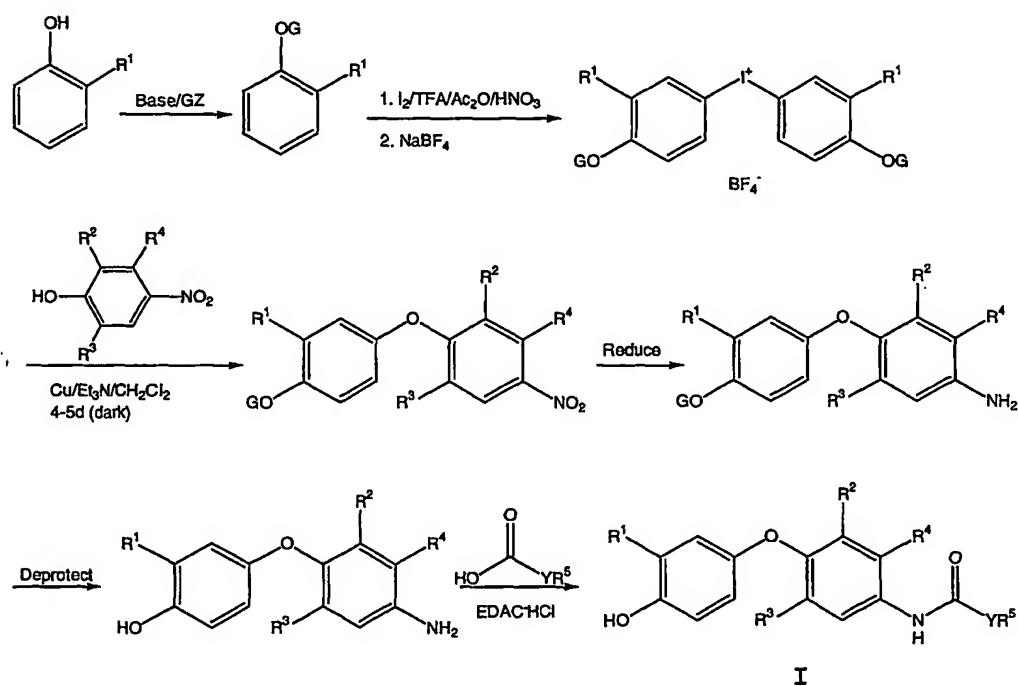
 If these indications are correct, then a TR β -selective agonist could be used to mimic a number of thyroid hormone actions, while having a lesser effect on the heart. Such a compound may be used for: (1)

25 replacement therapy in elderly subjects with hypothyroidism who are at risk for cardiovascular complications; (2) replacement therapy in elderly subjects with subclinical hypothyroidism who are at risk for cardiovascular complications; (3) obesity; (4)

30 hypercholesterolemia due to elevations of plasma LDL levels; (5) depression; and, (6) osteoporosis in combination with a bone resorption inhibitor.

Thyroid receptor ligands of the formula I, below, have previously been synthesized by several methods including the method summarized in Scheme A. (See United States Patent Application Serial No. 09/761,050, filed January 16, 2001.) The group G represents any group appropriate for protecting a hydroxyl moiety. The group Z represents a leaving group, such as a halogen. Examples of appropriate protecting groups G can be found, for example, in T. W. Greene & P. G. M. Wuts, "Protecting Groups in Organic Synthesis", 3rd Edition, Wiley, 1999.

Scheme A



15

This route is not optimal, however, due to the exothermicity of the formation of the iodonium triflate salt which is an unisolated intermediate in the synthesis depicted in Scheme A. In addition, the isolated, *bis*-phenyl iodonium salt is an intractable gummy solid at

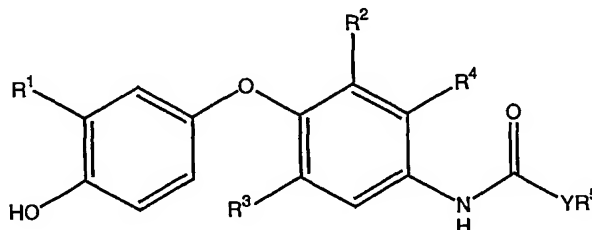
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room temperature, and the yield of the formation reaction is unpredictable. Further, the coupling reaction to form the mixed ether from the iodonium salt is not well adapted to commercial practice, because it takes place over an extended period of time, it requires unusual provisions to exclude light, and its yield is also unpredictable.

Accordingly, there is a need for improved processes for the preparation of thyroid receptor ligands, especially for processes that improve upon the safety and economic feasibility of the processes known in the art.

SUMMARY OF THE INVENTION

The present invention relates to the synthesis of compounds that have activity as thyroid receptor ligands, specifically compounds of formula I



20

wherein:

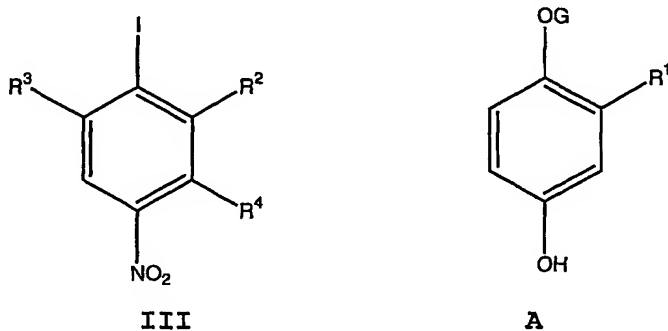
R¹ represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons;
R² and R³ are the same or different and represent
hydrogen, halogen, an alkyl group of 1 to 4 carbons, or a
cycloalkyl group of 3 to 6 carbons, at least one of R² and
R³ being other than hydrogen;
R⁴ represents hydrogen or a lower alkyl group;

R^5 represents a carboxylic acid or an alkyl ester thereof;
and

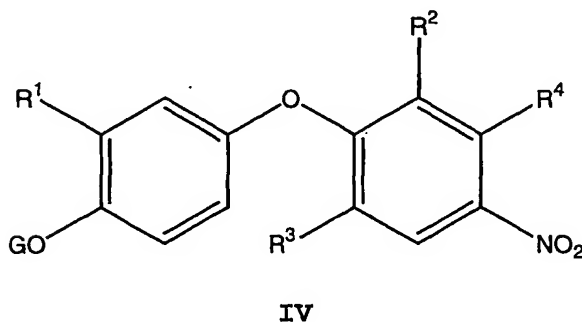
Y represents $-(CH_2)_n-$ where n is an integer from 1 to 5,
or a *cis*- or *trans*- ethylene group $-CH=CH-$;

5 and all stereoisomers thereof, prodrug esters thereof,
and pharmaceutically acceptable salts thereof.

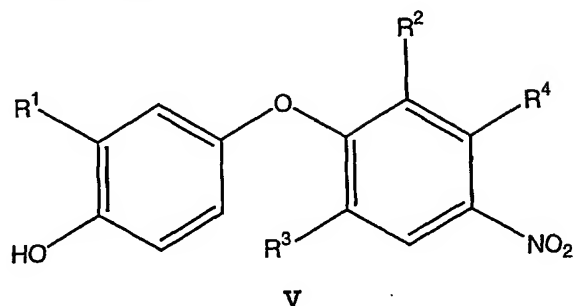
Compounds of formula I may be synthesized by
reacting a compound of formula III with a phenol of
10 formula A, wherein G is a protecting group, in the
presence of a base to produce a compound of formula IV,
wherein R^1 , R^2 , R^3 , and R^4 are as defined above;



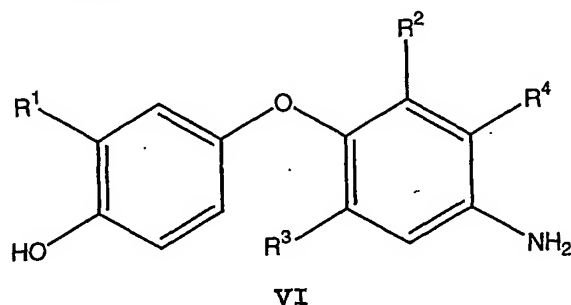
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deprotecting the compound of formula IV to form a compound of formula V;



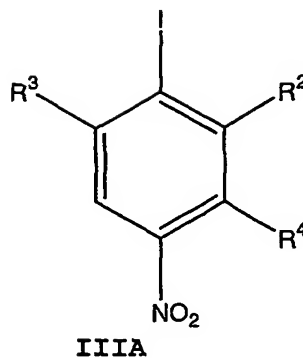
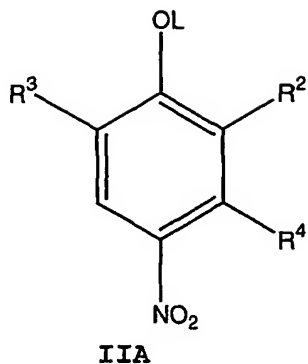
5 reducing the compound of formula V to produce a compound of formula VI;



10 reacting the compound of formula VI with a compound of the formula XC(O)YR^5 , wherein X represents OH or a halogen and R^5 is a carboxylic acid alkyl ester, in the presence of a base, to produce a compound of formula I wherein R^5 is a carboxylic acid alkyl ester; and

15 optionally de-esterifying the compound of formula I wherein R^5 is a carboxylic acid alkyl ester to produce a compound of formula I wherein R^5 is a carboxylic acid.

20 Also provided by the present invention are compounds that are novel starting materials of the above process having the formulae IIA and IIIA



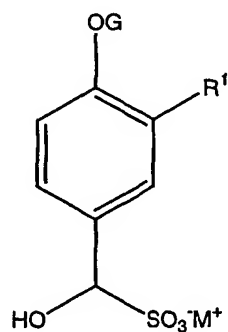
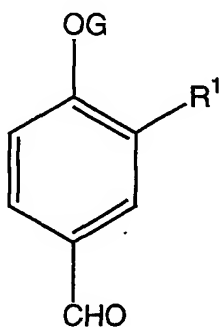
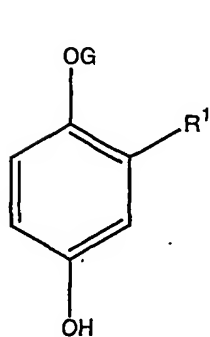
wherein

R^2 and R^3 independently represent bromo, chloro, or methyl;

R^4 represents hydrogen or methyl; and

L represents mesyl, tosyl, *p*-nitrobenzenesulfonyl, trihaloacetate, or triflate.

The present invention also provides compounds of the formulae A, A-I, and A-II



wherein M is a metal and G and R^1 are as defined above. In compounds of formula A, however, when G is a methyl group, R^1 is not a bromo, chloro, iodo, *t*-butyl, cyclohexyl, cyclopentyl, or isopropyl group.

Also provided by the present invention is a process for synthesizing a phenol of formula A, above, by reacting an aldehyde of formula A-I, above, with $M^+HSO_3^-$ wherein M is a metal to form a sulfonate of formula A-II, above. The sulfonate of formula A-II is reacted with an oxidant in the presence of a proton source to form the phenol of formula A.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph of the results of a temperature vs. time study of the formation reaction of the iodonium triflate salt of the comparative example.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The following definitions apply to the terms as used throughout this specification, unless otherwise limited in specific instances.

The term "thyroid receptor ligand" as used herein, is intended to cover any moiety which binds to a thyroid receptor. The ligand may act as an agonist, an antagonist, a partial agonist or a partial antagonist. Another term for "thyroid receptor ligand" is "thyromimetic".

Unless otherwise indicated, the term "lower alkyl", "alkyl" or "alk" as employed herein alone or as part of another group includes both straight and branched chain hydrocarbons, containing 1 to 12 carbons (in the case of alkyl or alk), in the normal chain, preferably 1 to 4 carbons, such as methyl, ethyl, propyl, isopropyl, butyl, t-butyl, or isobutyl, pentyl, hexyl, isohexyl, heptyl, 4,4-dimethylpentyl, octyl, 2,2,4-trimethylpentyl, nonyl,

decyl, undecyl, dodecyl. Each alkyl group may be optionally substituted with 1 to 4 substituents which may include alkyl, alkenyl, alkynyl, aryl, cycloalkyl, halogen, heteroaryl, hydroxy, cyano, nitro, amino and/or carboxyl or alkyl ester thereof.

The term "aryl" as employed herein alone or as part of another group refers to monocyclic and bicyclic aromatic groups containing 6 to 10 carbons in the ring portion (such as phenyl or naphthyl including 1-naphthyl and 2-naphthyl) and may be optionally substituted through available carbon atoms with 1, 2, or 3 groups selected from hydrogen, halo, alkyl, haloalkyl, alkoxy, haloalkoxy, alkenyl, trifluoromethyl, trifluoromethoxy, alkynyl, hydroxy, amino, nitro, cyano and/or carboxyl or an alkyl ester thereof.

The term "heteroaryl" or "heteroaromatic moiety" as used herein alone or as a part of another group refers to a 5- or 6-membered aromatic ring which includes 1, 2, 3, or 4 heteroatoms, one of which must be a nitrogen atom; the other heteroatoms when present may be nitrogen, oxygen or sulfur, and such rings may be fused to another aryl or heteroaryl ring. The heteroaryl groups may include one or more N-oxides. The heteroaryl group may optionally include 1 to 4 substituents such as aryl, alkyl, alkenyl, alkynyl, cycloalkyl, hydroxy, cyano, nitro, amino and/or carboxyl, or an alkyl ester thereof.

Unless otherwise indicated, the term "lower alkenyl" or "alkenyl" as used herein by itself or as part of another group refers to straight or branched chain radicals of 2 to 12 carbons, preferably 2 to 5 carbons,

in the normal chain, which include one to six double bonds in the normal chain, such as vinyl, 2-propenyl, 3-butenyl, 2-butenyl, 4-pentenyl, 3-pentenyl, 2-hexenyl, 3-hexenyl, 2-heptenyl, 3-heptenyl, 4-heptenyl, 3-octenyl, 3-nonenyl, 4-decenyl, 3-undecenyl, 4-dodecenyl, and the like, which may be substituted in the same manner as that described for alkyl groups.

Unless otherwise indicated, the term "lower alkynyl" or "alkynyl" as used herein by itself or as part of another group refers to straight or branched chain radicals of 2 to 12 carbons, preferably 2 to 8 carbons, in the normal chain; which include one triple bond in the normal chain, such as 2-propynyl, 3-butyne, 2-butyne, 4-pentyne, 3-pentyne, 2-hexyne, 3-hexyne, 2-heptyne, 3-heptyne, 4-heptyne, 3-octyne, 3-nonyne, 4-decyne, 3-undecyne, 4-dodecyne and the like, which may be substituted in the same manner as that described for alkyl groups.

Unless otherwise indicated, the term "cycloalkyl" as employed herein alone or as part of another group includes saturated cyclic hydrocarbon groups or partially unsaturated (containing 1 or 2 double bonds) cyclic hydrocarbon groups, containing one ring and a total of 3 to 7 carbons, preferably 3 to 6 carbons, forming the ring, which includes cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclopentenyl and cyclohexenyl, which may be substituted in the same manner as that described for alkyl groups.

The term "halogen" or "halo" as used herein alone or as part of another group refers to fluorine, chlorine,

bromine, and iodine, with chlorine or bromine being preferred.

The term "metal" as used herein refers to an element that can exist as a positive ion in solution. Suitable metals include alkali metals, transition metals, lanthanides, and actinides. Alkali metals are preferred. Lithium, sodium, and potassium are the most preferred metals.

As used herein, the term "about" means that amounts, sizes, formulations, parameters, and other quantities and characteristics are not and need not be exact, but may be approximate and/or larger or smaller, as desired, reflecting tolerances, conversion factors, rounding off, measurement error and the like, and other factors known to those of skill in the art. In general, an amount, size, formulation, parameter or other quantity or characteristic is "about" or "approximate" whether or not expressly stated to be such.

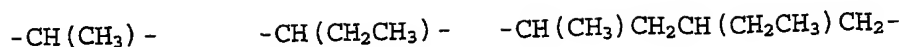
The divalent linking group represented by Y may be an alkylene group $-(CH_2)_n-$ that includes 1 to 5 carbons in the normal chain, and may also include 1, 2, or 3 alkyl substituents.

Examples of $(CH_2)_n$ groups include

$-CH_2-$ $-CH_2CH_2-$ $-CH_2CH_2CH_2-$ $-CH_2CH_2CH_2CH_2-$

$-CH_2CH_2C(CH_3)_2CH_2-$ $-CH_2CH_2CH(CH_2CH_3)CH_2-$

$-CH_2CH_2CH(CH_2CH_3)CH_2CH(CH_3)-$ $-C(CH_3)(CH_2CH_3)CH_2CH_2-$



Alternatively, Y may represent a *cis*- or *trans*-
5 alkylene group -CH=CH- which may be substituted as
described above.

The compounds of formula I may be present as salts,
in particular pharmaceutically acceptable salts. If the
10 compounds of formula I have, for example, at least one
basic center, they can form acid addition salts. These
are formed, for example, with strong inorganic acids,
such as mineral acids, for example sulfuric acid,
phosphoric acid or a hydrohalic acid, with strong organic
15 carboxylic acids, such as alkanecarboxylic acids of 1 to
4 carbon atoms which are unsubstituted or substituted,
for example, by halogen, for example acetic acid, such as
saturated or unsaturated dicarboxylic acids, for example
oxalic, malonic, succinic, maleic, fumaric, phthalic or
20 terephthalic acid, such as hydroxycarboxylic acids, for
example ascorbic, glycolic, lactic, malic, tartaric or
citric acid, such as amino acids, (for example aspartic
or glutamic acid or lysine or arginine), or benzoic acid,
or with organic sulfonic acids, such as (C1-C4) alkyl or
25 arylsulfonic acids which are unsubstituted or
substituted, for example by halogen, for example
methanesulfonic acid or *p*-toluenesulfonic acid.

Corresponding acid addition salts can also be formed
30 having, if desired, an additional basic center. The
compounds of formula I having at least one acid group
(for example COOH) can also form salts with bases.
Suitable salts with bases are, for example, metal salts,

such as alkali metal or alkaline earth metal salts, for example sodium, potassium or magnesium salts, or salts with ammonia or an organic amine, such as morpholine, thiomorpholine, piperidine, pyrrolidine, a mono-, di-, or tri-lower alkylamine, for example ethyl, *t*-butyl, diethyl, diisopropyl, triethyl, tributyl or dimethylpropylamine, or a mono, di, or trihydroxy lower alkylamine, for example mono, di or triethanolamine. Corresponding internal salts may furthermore be formed. Salts which are unsuitable for pharmaceutical uses but which can be employed, for example, for the isolation or purification of free compounds of formula I or their pharmaceutically acceptable salts, are also included.

Preferred salts of the compounds of formula I which include a basic group include monohydrochloride, hydrogensulfate, methanesulfonate, phosphate or nitrate.

Preferred salts of the compounds of formula I which include an acid group include sodium, potassium and magnesium salts and pharmaceutically acceptable organic amines.

Preferred compounds of formula I are those wherein Y represents $-(CH_2)_n-$ where *n* is 1 or 2; *R*¹ represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons; *R*² and *R*³ independently represent bromo, chloro, or methyl; *R*⁴ represents hydrogen or methyl; and *R*⁵ represents carboxyl.

More preferred compounds of formula I are those wherein

Y represents $-(CH_2)_n-$ where n is 1;

R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons, most preferred being isopropyl;

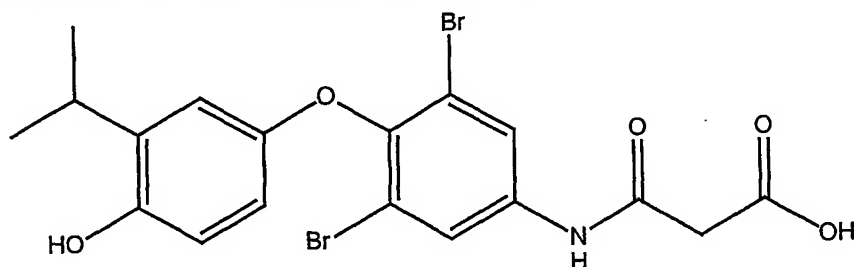
R^2 and R^3 independently represent bromo or chloro;

R^4 represents hydrogen or methyl; and

R^5 represents carboxyl.

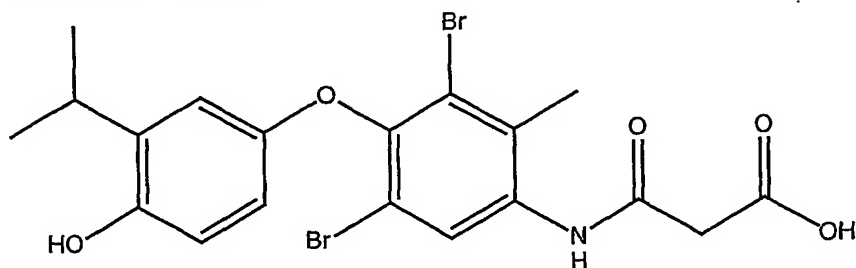
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Compounds of formula I that can be prepared by the methods of the invention include the following:

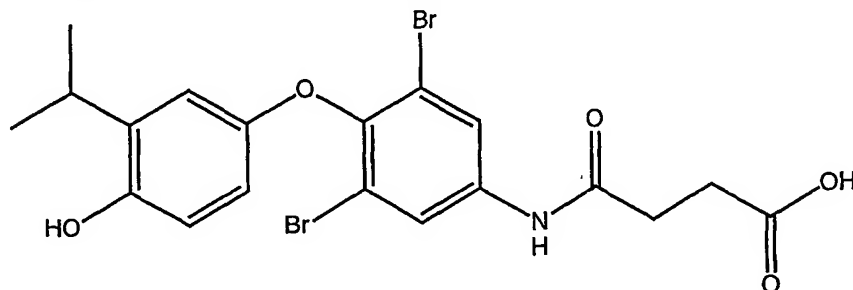


N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonic acid;

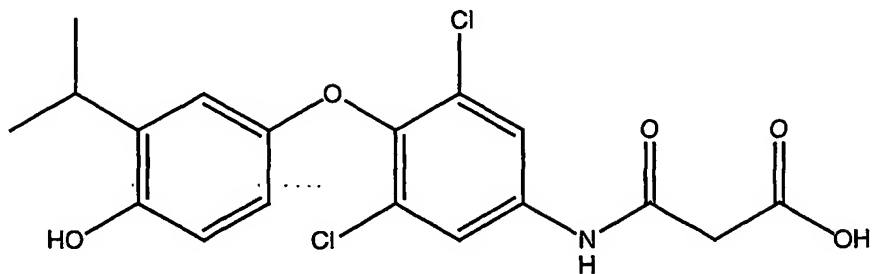
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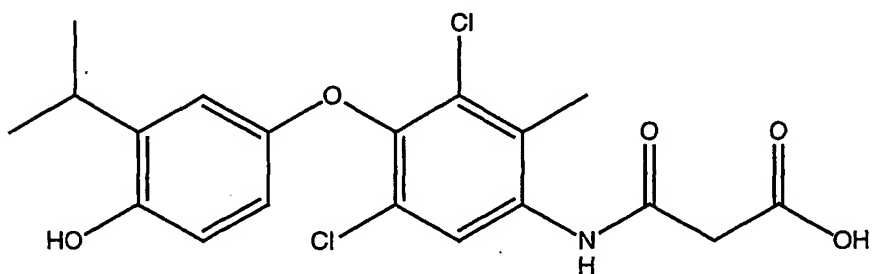
N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-2-methyl-phenyl]-malonic acid;



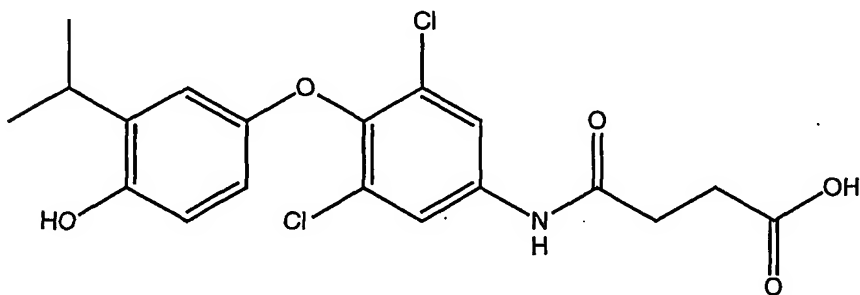
N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-succinamic acid;



5 *N*-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonamic acid;

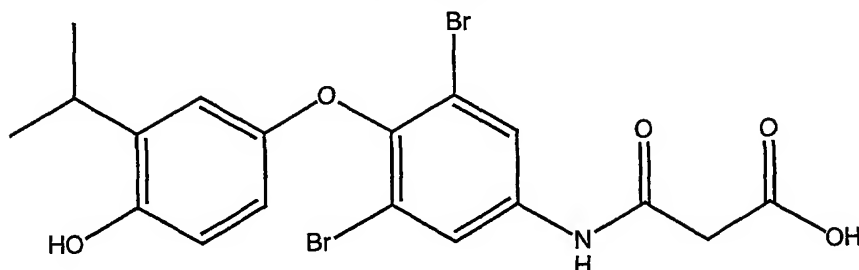


N-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-2-methyl-phenyl]-malonamic acid; and

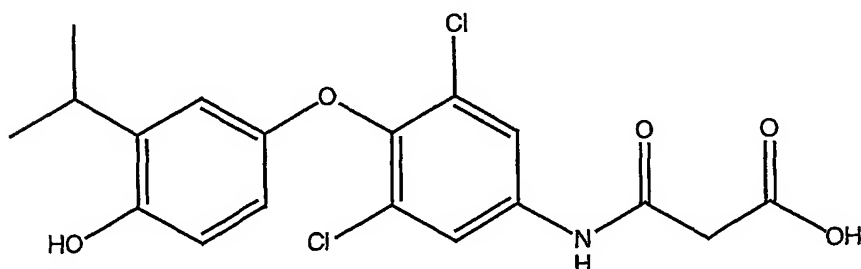


10 *N*-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-succinamic acid, or alkyl esters thereof, such as the methyl or ethyl ester.

The most preferred compounds of the invention are:



N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-
malonamic acid; and



N-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-
phenyl]-malonamic acid, or alkyl esters thereof, such as
the methyl or ethyl ester.

Preferred compounds of formula IIA include
trifluoro-methanesulfonic acid 2,6-dibromo-4-nitrophenyl
ester, trifluoro-methanesulfonic acid 2,6-dibromo-3-
methyl-4-nitrophenyl ester, trifluoro-methanesulfonic
acid 2,6-dichloro-4-nitrophenyl ester, and trifluoro-
methanesulfonic acid 2,6-dichloro-3-methyl-4-nitrophenyl
ester.

A more preferred compound of formula IIA is
trifluoro-methanesulfonic acid 2,6-dibromo-4-nitrophenyl
ester.

Preferred compounds of formula IIIA include 1,3-dibromo-2-iodo-5-nitrobenzene, 1,3-dibromo-2-iodo-4-methyl-5-nitrobenzene, 1,3-dichloro-2-iodo-5-nitrobenzene, and 1,3-dichloro-2-iodo-4-methyl-5-nitrobenzene.

A more preferred compound of formula IIIA is 1,3-dibromo-2-iodo-5-nitrobenzene.

Preferred compounds of formulae A, A-I, and A-II for practicing the present invention are those in which R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons.

More preferred compounds of formulae A, A-I, and A-II are those in which R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons, most preferred being isopropyl.

The most preferred compound of formula A-I is 3-isopropyl-4-methoxybenzaldehyde. The most preferred compound of formula A-II is hydroxy-(3-isopropyl-4-methoxyphenyl)methanesulfonic acid, sodium salt. The most preferred phenol of formula A is 3-isopropyl-4-methoxyphenol.

All stereoisomers of the compounds of formula I are contemplated, either in admixture or in pure or substantially pure form. The compounds of formula I can have asymmetric centers at any of the carbon atoms including any one of the R substituents. Consequently, compounds of formula I can exist in enantiomeric or

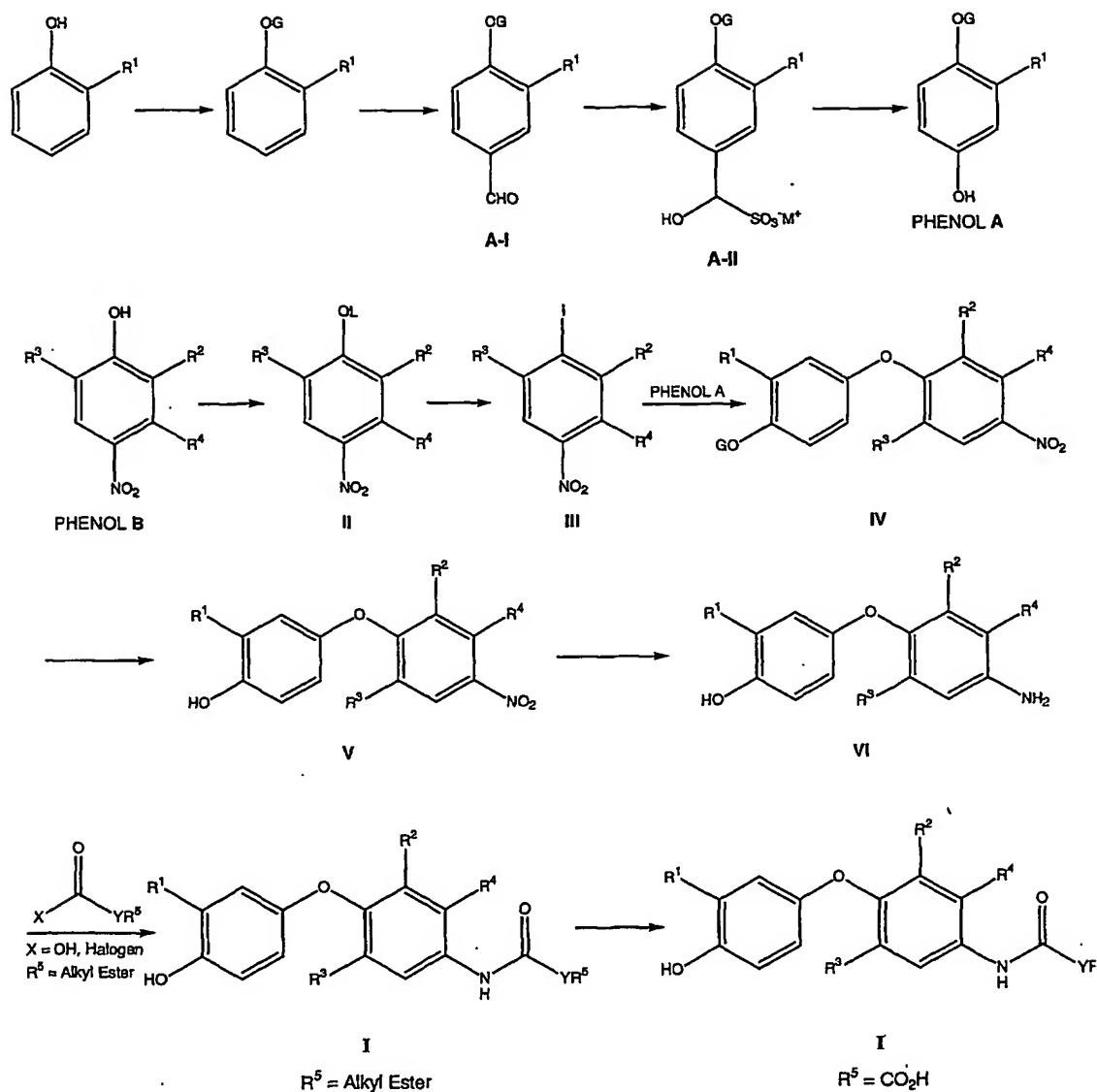
diastereomeric forms or in mixtures thereof. The processes for preparation can utilize racemates, enantiomers or diastereomers as starting materials. When diastereomeric or enantiomeric products are prepared, they can be separated by conventional methods, for example, chromatographic or fractional crystallization.

The compounds of formula I may be prepared by the exemplary processes described in the following reaction schemes. Exemplary reagents and procedures for these reactions appear hereinafter and in the working examples. Protection and deprotection in the Schemes herein may be carried out by procedures generally known in the art (see, for example, Greene & Wuts).

15

Scheme 1 sets forth an improved method of synthesizing compounds of formula I.

SCHEME 1



5

In scheme 1, R^1 , R^2 , R^3 , and R^4 are as defined above, M represents a metal, the group G represents any group appropriate for protecting a hydroxyl moiety, and the group -OL represents a leaving group. Examples of suitable protecting groups G can be found in Green and Wuts, for example. Examples of suitable leaving groups L can be found in Carey, F.A. and Sundberg, R.J., *Advanced*

10

Organic Chemistry (1977, Plenum Press, New York), Part A, Section 5.6, for example.

Preferred protecting groups include, e.g.,
5 substituted or unsubstituted alkyl, methoxymethyl (MOM),
carboxybenzyl (CBz), benzyloxymethyl (BOM), 2-
(trimethylsilyl) ethoxymethyl (SEM), aryl thiomethyl
(MTM), tetrahydropyranyl, methyl 2-propenyl ether, ethyl
vinyl ether, triphenylmethyl (trityl) and other
10 triarylmethyl groups, esters, carbonates, phosphinates,
sulfonates, and silyl groups.

More preferred protecting groups are alkyl groups.

15 The most preferred protecting groups are alkyl
groups containing 1 to 6 carbons, especially methyl
groups.

Preferred leaving groups include, e.g., tri-
20 haloacetates and sulfonate esters.

More preferred leaving groups are trihalo-
methylsulfonate, methylsulfonate (mesyl), *p*-
nitrobenzenesulfonate, *p*-toluenesulfonate (tosyl), and
25 trihaloacetate groups.

The most preferred leaving groups are
trifluoromethylsulfonate (triflate) groups.

30 Phenols of formula A may be synthesized by methods
known in the art. See, for example, Blank, B., *J. Med*
Chem., 1963, 6, 554, and Bowman et al., *J. Chem. Soc. C*,
1966, 2274. Phenols of formula A may also be synthesized

by the novel method of the present invention, wherein an aldehyde of formula A-I is reacted with about 2 to 4 equivalents of a sulfite reagent, such as an sodium sulfite or an alkali hydrogen sulfite, at about 20° to 30°C to produce a sulfonate of formula A-II. Preferably, about 2.25 to 2.75 equivalents of sulfite reagent are used, the preferred sulfite reagent being sodium sulfite.

The phenol of formula A is formed by reacting the sulfonate of formula A-II with an oxidant, in the presence of a proton source. Preferably, about 1 to 2 equivalents of protons and about 3 to 5 equivalents of oxidant are used.

Suitable oxidants for use in the synthesis of Phenol A include, e.g., peroxybenzoic acids, alkali metal monoperoxophthalates, hydrogen peroxide optionally catalyzed by Me-ReO_3 , oxygen in the presence of a copper species, H_3PO_5 , and oxone. More preferably, the oxidant is hydrogen peroxide.

Preferably, the proton source is a protic acid. More preferably, the proton source is *p*-toluenesulfonic acid.

25

Phenol B may be synthesized by well-known methods from commercially available starting materials. See, for example, Yenes, S. et al., *Tetrahedron*, 1999, 55, 14111; Bozell, J. et al., *Tet. Lett.*, 1998, 39, 2261; Fischer, A. et al., *Tet. Lett.*, 1988, 29, 1869; Rodygin, M. et al., *Zh. Org. Khim.*, 1992, 28, 1926; Xu, F. et al., *Shanghai. Keji Daxue Xuebao*, 1989, 12, 72; Clewly, R. et al., *Can. J. Chem.*, 1989, 67, 1472; Kajigaeshi, S. et

30

al., *Bull. Chem. Soc. Japan*, 1987, 60, 4187; Kajigaeshi
S. et al., *Chem. Lett.*, 1987, 4, 627; Lemaire, M. et al.,
Tetrahedron, 1987, 43, 835; Fujita, S. et al., JP
03291241, 1991; Clewly, R. et al., *Tetrahedron*, 1989, 45,
1299; Kajigaeshi, S. et al., *Chem. Express*, 1990, 5, 141;
Kakinami, T. et al., *Bull. Chem. Soc. Japan*, 1989, 62,
3373; Kajigaeshi, S. et al., *Technol. Rep. Yamaguchi
Univ.*, 1987, 4, 65; Gaude D. et al., *Can. J. Chem.*, 1989,
67, 104; Kajigaeshi, S. et al., *Chem. Lett.*, 1987, 2109.
Certain Phenol B compounds are also commercially
available from the Aldrich Company, for example.

Compounds of formula III are synthesized from
compounds of formula II by iodination. Iodination
reagents include, e.g., alkali metal iodides and
tetraalkylammonium iodides. Preferred iodination
reagents include, e.g., sodium iodide and potassium
iodide. Preferably, the iodination agent is present in
about 1-fold to 10-fold excess. Preferably, the
iodination is carried out at 80° to 110°C. Preferably,
the solvent is acetone, methyl ethyl ketone (MEK), methyl
isobutyl ketone (MIBK), toluene,
hexamethylphosphorotriamide (HMPA), dimethyl formamide
(DMF), or a higher-boiling acetate. More preferably, the
iodination is carried out by reacting the compound of
formula II with 3- to 5-fold excess NaI or KI in DMF, MEK
or MIBK at a temperature of about 75° to 100°C.

The coupling reaction to form the mixed ether of
formula IV from the compound of formula III and phenol A
is conducted in the presence of a base. Preferred bases
include, e.g., alkyl lithiums, lithium diisopropylamine,
Grignard reagents, alkali metal hydrides, alkali metal

carbonates, alkali metal hydroxides, and alkali metal alkoxides. Preferred solvents include THF, 1,2-dimethoxyethane, 1,1-dimethoxyethane, dimethyl sulfoxide (DMSO), and toluene. Preferably, about 1.1 to 1.5
5 equivalents of base are used. Preferably, the reaction temperature is between about -78° and 30°C. More preferably, the coupling reaction is performed in toluene/DMF or toluene/THF or DMF at room temperature using potassium hexamethyldisilazide (KHMDs), potassium
10 t-butoxide, potassium t-pentoxide, LiHMDS, or NaHMDS as the base.

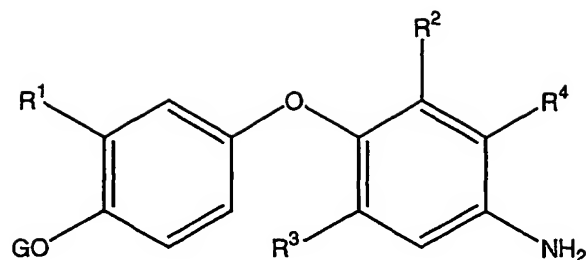
The deprotection of the protected hydroxyl group of compound IV may be carried out at reaction temperatures
15 below room temperature. Preferred deprotecting agents include, e.g., trialkylsilyl halides, mercaptans, boron halides, magnesium halides, aluminum halides, hydrogen over palladium catalyst, or iron halides. Preferably, the reaction temperature is maintained between about 0°
20 to 15°C. Preferably, the deprotection reaction is quenched with at least about 4 molar equivalents of methanol. More preferably, the deprotecting agent is a trialkylsilyl halide or BBr₃.

25 Reducing agents and conditions suitable for use in reducing the formula V compound include, e.g., elemental metals such as Fe and Na; hydrogen catalyzed by nickel, palladium or platinum; alkali metal borohydrides; magnesium-mediated reductions; iron carbonyls; iron
30 sulfate mediated reductions; iron and HCl; sodium hydrogen sulfite; sodium dithionite; tin and tin halide mediated reductions; and alkali metal sulfides and polysulfides. Preferred reducing agents include, e.g.,

sodium hydrogen sulfite, sodium dithionite, or iron and HCl. Preferably, about 4 to 6 equivalents of reducing agent are used. Preferably, the reaction temperature is maintained between about 25° to 50°C. More preferably, the reducing agent is sodium dithionite, and the reaction is carried out over about 1h in THF/H₂O at about 50°C.

Formula VI compounds may be converted into formula I compounds wherein R⁵ is an alkyl ester group by reaction with a compound of formula XC(O)YR⁵, wherein X and Y are as defined above, under basic conditions. Preferred bases are alkali and alkaline earth metal hydroxides and alkoxides, alkyl amines, and substituted alkyl amines. Preferred compounds of formula XC(O)YR⁵ are alkyl and malonyl halides. Preferably, the reaction temperature is maintained between about 0° and 5°C. Preferably, about 1 equivalent or a slight excess of base is used. Preferably, about 50% to 100% excess of the compound of formula XC(O)YR⁵ is used. More preferably, XC(O)YR⁵ is ethyl or methyl malonyl chloride and the reaction is carried out at a temperature around 0°C under basic conditions, using NaOH in t-butyl methyl ether (TBME).

In the above Scheme 1, the deprotection of the phenol hydroxyl group of compound IV to form the compound of formula V is depicted as being performed before the reduction of the nitro group to form the compound of formula VI. The order in which these two reactions are performed may be reversed, however. When the order is thus reversed, a compound of formula VA is formed by the reduction.



VA

The compound of formula VA is readily deprotected to form
 5 the compound of formula VI. It is preferred, however, to
 carry out the deprotection before the reduction.

Compounds of formula I wherein R⁵ is an alkyl ester
 may be de-esterified by hydrolysis under acidic or basic
 10 conditions to form compounds of formula I wherein R⁵ is a
 carboxylic acid. Examples of suitable de-esterifying
 agents include, e.g., enzymes; hydrogen over catalytic
 palladium; lithium and ammonia; zinc; alkali metal
 hydroxides and alkoxides; alkaline earth metal hydroxides
 15 and alkoxides; sodium sulfide; KSCN; thioethers in
 combination with Lewis acids; anions of alkyl thiols or
 selenols; trialkyl silyl halides; zinc dichloride;
 mercury (II) acetate; and strong protic acids.

Preferred hydrolysis reagents and conditions
 20 include, e.g., an alkali metal hydroxide in the presence
 of an ether such as tetrahydrofuran or *tert*-butyl methyl
 ether. Most preferably, the hydrolysis is carried out *in*
situ by the addition of approximately 1.5 to 2 molar
 25 equivalents of NaOH to the reaction mixture at a
 temperature around 0°C after the reaction of compound VI
 with XC(O)YR⁵ wherein X and Y are as previously defined

and R⁵ is an alkyl ester group is observed to be essentially complete.

5 The compounds made by the processes of the invention are agonists that are preferably selective for the thyroid hormone receptor-beta, and as such are useful in the treatment of obesity, hypercholesterolemia and atherosclerosis by lowering of serum LDL levels, alone or optionally in combination with a lipid modulating drug
10 such as an HMG CoA reductase inhibitor, fibrate, MTP inhibitor, squalene synthetase inhibitor and/or other hypolipidemic agent and/or optionally in combination with an antidiabetic agent; useful in the amelioration of depression, alone or optionally in combination with an
15 antidepressant such as fluoxetine and desipramine; and useful in the stimulation of bone formation to treat osteoporosis, alone or optionally in combination with any known bone resorption inhibitor such as alendronate sodium.

20

 In addition, the compounds of the invention may be useful as replacement therapy in elderly patients with hypothyroidism or subclinical hypothyroidism who are at risk for cardiovascular complications, in the treatment
25 of the elderly to provide a sense of well-being, and in the treatment of non-toxic goiter; in the management of papillary or follicular thyroid cancer (alone or with T₄); in the treatment of skin disorders such as psoriasis; and in the treatment of glaucoma,
30 cardiovascular disease such as in the prevention or treatment of atherosclerosis, and congestive heart failure.

The compounds made by the processes of the invention may be employed alone or in combination with an appetite suppressant such as sibutramine, and/or in combination with anti-obesity agents such as orlistat, and/or in
5 combination with a $\beta 3$ agonist, for treating obesity.

The compounds made by the processes of the invention may also be used to treat skin disorders or diseases involving dermal atrophy such as glucocorticoid induced
10 dermal atrophy, including restoration of dermal atrophy induced by topical glucocorticoids, the prevention of dermal atrophy induced by topical glucocorticoids (such as the simultaneous treatment with topical glucocorticoid or a pharmacological product including both
15 glucocorticoid and a compound of the invention), the restoration/prevention of dermal atrophy induced by systemic treatment with glucocorticoids, restoration/prevention of atrophy in the respiratory system induced by local treatment with glucocorticoids,
20 UV-induced dermal atrophy, or dermal atrophy induced by aging (wrinkles, etc.), wound healing, keloids, stria, cellulite, roughened skin, actinic skin damage, lichen planus, ichthyosis, acne, psoriasis, Dernier's disease, eczema, atopic dermatitis, chloracne, pityriasis and skin
25 scarring.

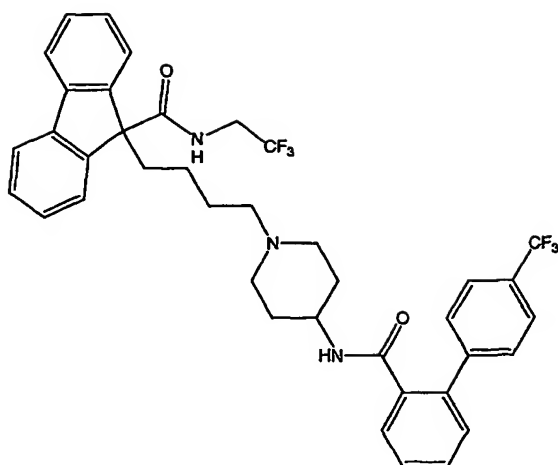
In treating skin disorders or diseases as described above, the compounds made by the processes of the invention may be used alone or optionally in combination
30 with a retinoid such as tretinoin or a vitamin D analog, employing amounts as disclosed in the PDR.

The hypolipidemic agent which may be optionally employed in combination with the compounds of formula I may include thiazolidinediones, MTP inhibitors, HMG CoA reductase inhibitors, squalene synthetase inhibitors, fibric acid derivatives, ACAT inhibitors, cholesterol absorption inhibitors, ileal Na⁺/bile acid cotransporter inhibitors, bile acid sequestrants, and/or nicotinic acid and derivatives thereof.

MTP inhibitors employed herein include MTP inhibitors disclosed in U.S. Patent No. 5,595,872, U.S. Patent No. 5,739,135, U.S. Patent No. 5,712,279, U.S. Patent No. 5,760,246, U.S. Patent No. 5,827,875, U.S. Patent No. 5,885,983 and U.S. Patent No. 5,962,440. Preferred are each of the preferred MTP inhibitors disclosed in each of the above patents and applications.

Most preferred MTP inhibitors to be employed with the compounds made by the processes of the present invention include preferred MTP inhibitors as set out in U.S. Patent Nos. 5,739,135 and 5,712,279, and U.S. Patent No. 5,760,246.

The most preferred MTP inhibitor is 9-[4-[4-[[2-(2,2,2-Trifluoroethoxy)benzoyl]amino]-1-piperidinyl]butyl]-N-(2,2,2-trifluoroethyl)-9H-fluorene-9-carboxamide, formula VII.



VII

5 The hypolipidemic agent may be an HMG CoA reductase inhibitor which includes, but is not limited to, mevastatin and related compounds as disclosed in U.S. Patent No. 3,983,140, lovastatin (mevinolin) and related compounds as disclosed in U.S. Patent No. 4,231,938, pravastatin and related compounds such as disclosed in 10 U.S. Patent No. 4,346,227, simvastatin and related compounds as disclosed in U.S. Patent Nos. 4,448,784 and 4,450,171. Other HMG CoA reductase inhibitors which may be employed herein include, but are not limited to, 15 fluvastatin, disclosed in U.S. Patent No. 5,354,772, cerivastatin disclosed in U.S. Patent Nos. 5,006,530 and 5,177,080, atorvastatin disclosed in U.S. Patent Nos. 4,681,893, 5,273,995, 5,385,929 and 5,686,104, pyrazole analogs of mevalonolactone derivatives as disclosed in 20 U.S. Patent No. 4,613,610, indene analogs of mevalonolactone derivatives as disclosed in PCT application WO 86/03488, 6-[2-(substituted-pyrrol-1-yl)-alkyl]pyran-2-ones and derivatives thereof as disclosed in U.S. Patent No. 4,647,576, Searle's SC-45355 (a 3-

substituted pentanedioic acid derivative)
dichloroacetate, imidazole analogs of mevalonolactone as
disclosed in PCT application WO 86/07054, 3-carboxy-2-
hydroxy-propane-phosphonic acid derivatives as disclosed
5 in French Patent No. 2,596,393, 2,3-disubstituted
pyrrole, furan and thiophene derivatives as disclosed in
European Patent Application No. 0221025, naphthyl analogs
of mevalonolactone as disclosed in U.S. Patent No.
4,686,237, octahydronaphthalenes such as disclosed in
10 U.S. Patent No. 4,499,289, keto analogs of mevinolin
(lovastatin) as disclosed in European Patent Application
No. 0,142,146 A2, as well as other known HMG CoA
reductase inhibitors.

15 In addition, phosphinic acid compounds useful in
inhibiting HMG CoA reductase suitable for use with
compounds of formula I are disclosed in GB 2205837.

The squalene synthetase inhibitors suitable for use
20 with compounds of formula I include, but are not limited
to, α -phosphonosulfonates disclosed in U.S. Patent No.
5,712,396, those disclosed by Biller et al., J. Med.
Chem., 1988, Vol. 31, No. 10, pp 1869-1871, including
isoprenoid (phosphinylmethyl)phosphonates as well as
25 other squalene synthetase inhibitors as disclosed in U.S.
Patent Nos. 4,871,721 and 4,924 024 and in Biller et al.,
Current Pharmaceutical Design, 2, 1-40 (1996).

In addition, other squalene synthetase inhibitors
30 suitable for use with compounds of formula I include the
terpenoid pyrophosphates disclosed by P. Ortiz de
Montellano et al., J. Med. Chem., 1977, 20, 243-249, the
farnesyl diphosphate analog A and presqualene

pyrophosphate (PSQ-PP) analogs as disclosed by Corey and Volante, J. Am. Chem. Soc., 1976, 98, 1291-1293, phosphinylphosphonates reported by McClard, R.W. et al, J. Am. Chem. Soc., 1987, 109, 5544 and cyclopropanes reported by Capson, T.L., PhD dissertation, June, 1987, Dept. Med. Chem. Univ. of Utah, Abstract, Table of Contents, pp 16, 17, 40-43, 48-51, Summary.

Other hypolipidemic agents suitable for use with compounds of formula I include, but are not limited to, fibric acid derivatives, such as fenofibrate, gemfibrozil, clofibrate, bezafibrate, ciprofibrate, clinofibrate and the like, probucol, and related compounds as disclosed in U.S. Patent No. 3,674,836, probucol and gemfibrozil being preferred, bile acid sequestrants such as cholestyramine, colestipal and DEAE-Sephadex (Secholex™, Policexide™, as well as lipostabil (Rhone-Poulenc), Eisai E-5050 (an N-substituted ethanolamine derivative), imanixil (HOE-402), tetrahydrolipstatin (THL), istigmastanylphosphorylcholine (SPC, Roche), aminocyclodextrin (Tanabe Seiyoku), Ajinomoto AJ-814 (azulene derivative), melinamide (Sumitomo), Sandoz 58-035, American Cyanamid CL-277,082 and CL-283,546 (disubstituted urea derivatives), nicotinic acid, acipimox, acifran, neomycin, p-aminosalicylic acid, aspirin, poly(diallylmethylamine) derivatives such as disclosed in U.S. Patent No. 4,759,923, quaternary amine poly(diallyldimethylammonium chloride) and ionenes such as disclosed in U.S. Patent No. 4,027,009, and other known serum cholesterol lowering agents.

The other hypolipidemic agent may be an ACAT inhibitor such as disclosed in Drugs of the Future 24, 9-15 (1999), (Avasimibe); Nicolosi et al., Atherosclerosis (Shannon, Ireland) 1998, 137(1), 77-85; Ghiselli, 5 Giancarlo, Cardiovasc. Drug Rev. 1998, 16(1), 16-30; Smith, C., et al., Bioorg. Med. Chem. Lett. 1996, 6(1), 47- 50; Krause et al., Editor(s): Ruffolo et al., Inflammation: Mediators Pathways 1995, 173-98, Publisher: CRC Press, Boca Raton, Fla.; Sliskovic et al., Curr. Med. 10 Chem. 1994, 1(3), 204-25; Stout et al., Chemtracts: Org. Chem. 1995, 8(6), 359-62.

The hypolipidemic agent may be a cholesterol absorption inhibitor preferably Schering-Plough's 15 SCH48461 as well as those disclosed in Atherosclerosis 115, 45-63 (1995) and J. Med. Chem. 41, 973 (1998).

The hypolipidemic agent may be an ileal Na⁺/bile acid cotransporter inhibitor such as disclosed in Drugs 20 of the Future, 24, 425-430 (1999). Preferred hypolipidemic agents are pravastatin, lovastatin, simvastatin, atorvastatin, fluvastatin and cerivastatin.

The amounts and dosages employed will be as 25 indicated in the Physician's Desk Reference (PDR) and/or in the patents set out above.

The compounds of formula I will be employed in a weight ratio to the hypolipidemic agent, the 30 antidepressant, and/or bone resorption inhibitor and/or appetite suppressant (where present), within the range from about 500:1 to about 0.005:1, preferably from about 300:1 to about 0.01:1.

The antidiabetic agent which may be optionally employed in combination with compounds of formula I may include biguanides, sulfonyl ureas, glucosidase inhibitors, thiazolidinediones and/or α P2 inhibitors and/or PPAR α agonists, PPAR γ agonists or PPAR α/γ dual agonists, and/or SGLT2 inhibitors, or meglitinide.

The antidiabetic agent may be an oral antihyperglycemic agent, preferably a biguanide such as metformin or phenformin or salts thereof.

Where the antidiabetic agent is a biguanide, the compounds of formula I will be employed in a weight ratio to biguanide within the range from about 0.01:1 to about 100:1, preferably from about 0.5:1 to about 2:1.

The antidiabetic agent may also preferably be a sulfonylurea such as glyburide (also known as glibenclamide), glimepiride (disclosed in U.S. Patent No. 4,379,785), glipizide, gliclazide or chlorpropamide, other known sulfonylureas or other antihyperglycemic agents which act on the ATP-dependent channel of the β -cells, with glyburide and glipizide being preferred.

The compounds of formula I will be employed in a weight ratio to the sulfonyl urea in the range from about 0.01:1 to about 100:1, preferably from about 0.2:1 to about 10:1.

The oral antidiabetic agent may also be a glucosidase inhibitor such as acarbose (disclosed in U.S. Patent No. 4,904,769) or miglitol (disclosed in

U.S. Patent No. 4,639,436), which may be administered in a separate oral dosage form.

5 The compounds of formula I will be employed in a weight ratio to the glucosidase inhibitor within the range from about 0.01:1 to about 100:1, preferably from about 0.5:1 to about 50:1.

10 The compounds of formula I may be employed in combination with a thiazolidinedione oral anti-diabetic agent or other insulin sensitizers (which has an insulin sensitivity effect in NIDDM patients) such as troglitazone (Warner-Lambert's Rezulin™, disclosed in U.S. Patent No. 4,572,912), rosiglitazone (SKB),
15 pioglitazone (Takeda), Mitsubishi's MCC-555 (disclosed in U.S. Patent No. 5,594,016), Glaxo-Welcome's GI-262570, englitazone (CP-68722, Pfizer), or darglitazone (CP-86325, Pfizer).

20 The compounds of formula I will be employed in a weight ratio to the thiazolidinedione in an amount within the range from about 0.01:1 to about 100:1, preferably from about 0.5:1 to about 5:1.

25 The sulfonylurea and thiazolidinedione in amounts of less than about 150 mg oral antidiabetic agent may be incorporated in a single tablet with the compounds of structure I.

30 The compounds of formula I may also be employed in combination with a non-oral antihyperglycemic agent such as insulin or with glucagon-like peptide-1 (GLP-1) such as GLP-1(1-36) amide, GLP-1(7-36) amide, GLP-1(7-37) (as

disclosed in U.S. Patent No. 5,614,492 to Habener, the disclosure of which is incorporated herein by reference), which may be administered via injection, intranasal, or by transdermal or buccal devices.

5

Where present, metformin, the sulfonylureas, such as glyburide, glimepiride, glipyrizide, glipizide, chlorpropamide and gliclazide and the glucosidase inhibitors acarbose or miglitol or insulin may be employed in formulations as described above and in amounts and dosing as indicated in the Physician's Desk Reference (PDR).

15

Where present, metformin or salt thereof may be employed in amounts within the range from about 500 to about 2000 mg per day which may be administered in single or divided doses one to four times daily.

20

Where present, the thiazolidinedione anti-diabetic agent may be employed in amounts within the range from about 0.01 to about 2000 mg/day, which may be administered in single or divided doses one to four times per day.

25

Where present insulin may be employed in formulations, amounts and dosing as indicated by the Physician's Desk Reference (PDR).

30

Where present GLP-1 peptides may be administered in oral buccal formulations, by nasal administration or parenterally as described in U.S. Patent Nos. 5,346,701 (TheraTech), 5,614,492 and 5,631,224.

The antidiabetic agent may also be a PPAR α/γ dual agonist such as disclosed by Murakami et al., "A Novel Insulin Sensitizer Acts As a Coligand for Peroxisome Proliferation -Activated Receptor Alpha (PPAR alpha) and PPAR gamma. Effect on PPAR alpha Activation on Abnormal Lipid Metabolism in Liver of Zucker Fatty Rats", Diabetes 47, 1841-1847 (1998).

The antidiabetic agent may be an α P2 inhibitor such as disclosed in U.S. application Serial No. 09/391,053, filed September 7, 1999, and U.S. provisional application No. 60/127,745, filed April 5, 1999, employing dosages as set out herein.

The antidiabetic agent may be an SGLT2 inhibitor such as disclosed in U.S. provisional application 60/158,773, filed October 12, 1999.

The compounds of formula I will be employed in a weight ratio to the PPAR α agonist, PPAR γ agonist, PPAR α/γ dual agonists, SGLT2 inhibitor and/or α P2 inhibitor within the range from about 0.01:1 to about 100:1, preferably from about 0.5:1 to about 5:1.

The dose administered must be carefully adjusted according to age, weight and condition of the patient, as well as the route of administration, dosage form and regimen and the desired result.

The dosages and formulations for the hypolipidemic agent and antidiabetic agent will be as disclosed in the various patents and applications discussed above and in the Physicians' Desk Reference.

The dosages and formulations for the other hypolipidemic agent, antidepressant, bone resorption inhibitor, appetite suppressant and anti-obesity agent to be employed, where applicable, will be as set out in the latest edition of the Physicians' Desk Reference.

For oral administration, a satisfactory result may be obtained employing the MTP inhibitor in an amount within the range of from about 0.01 mg/kg to about 100 mg/kg and preferably from about 0.1 mg/kg to about 75 mg/kg, one to four times daily.

A preferred oral dosage form, such as tablets or capsules, will contain the MTP inhibitor in an amount of from about 1 to about 500 mg, preferably from about 2 to about 400 mg, and more preferably from about 5 to about 250 mg, one to four times daily.

For parenteral administration, the MTP inhibitor will be employed in an amount within the range of from about 0.005 mg/kg to about 10 mg/kg and preferably from about 0.005 mg/kg to about 8 mg/kg, one to four times daily.

For oral administration, a satisfactory result may be obtained employing an HMG CoA reductase inhibitor, for example, pravastatin, lovastatin, simvastatin, atorvastatin, fluvastatin or cerivastatin in dosages employed as indicated in the Physician's Desk Reference, such as in an amount within the range of from about 1 to 2000 mg, and preferably from about 4 to about 200 mg.

The squalene synthetase inhibitor may be employed in dosages in an amount within the range of from about 10 mg to about 2000 mg and preferably from about 25 mg to about 200 mg.

5

A preferred oral dosage form, such as tablets or capsules, will contain the HMG CoA reductase inhibitor in an amount from about 0.1 to about 100 mg, preferably from about 5 to about 80 mg, and more preferably from about 10 to about 40 mg.

10

A preferred oral dosage form, such as tablets or capsules will contain the squalene synthetase inhibitor in an amount of from about 10 to about 500 mg, preferably from about 25 to about 200 mg.

15

The compounds of formula I and the hypolipidemic agent, antidepressant or bone resorption inhibitor may be employed together in the same oral dosage form or in separate oral dosage forms taken at the same time.

20

The compositions described above may be administered in the dosage forms as described above in single or divided doses of one to four times daily. It may be advisable to start a patient on a low dose combination and work up gradually to a high dose combination.

25

The preferred hypolipidemic agent is pravastatin, simvastatin, lovastatin, atorvastatin, fluvastatin or cerivastatin.

30

The compounds of formula I can be administered orally or parenterally such as subcutaneously or

intravenously, as well as by nasal application, rectally or sublingually to various mammalian species known to be subject to such maladies, e.g., humans, cats, dogs and the like in an effective amount within the dosage range
5 of about 0.1 to about 100 mg/kg, preferably about 0.2 to about 50 mg/kg and more preferably about 0.5 to about 25 mg/kg (or from about 1 to about 2500 mg, preferably from about 5 to about 2000 mg) on a regimen in single or 2 to 4 divided daily doses.

10

The active substance can be utilized in a composition such as tablet, capsule, ointment, hydrophilic ointment, cream, lotion, solution or suspension or in other type carrier materials such as
15 transdermal devices, iontophoretic devices, rectal suppositories, inhalant devices and the like. The composition or carrier will contain about 5 to about 500 mg per unit of dosage of a compound of formula I. The active substances may be compounded in a conventional
20 matter with a physiologically acceptable vehicle or carrier, excipient, binder, preservative, stabilizer, or flavor, as called for by accepted pharmaceutical practice.

25

The following examples are provided to describe the invention in further detail. These examples are intended to illustrate and not to limit the invention. All temperatures are given in centigrade degrees (°C) unless otherwise noted.

30

EXAMPLE 1**Preparation of 1-isopropyl-2-methoxybenzene**

A 22L three-necked round bottom flask equipped with mechanical stirring was charged with KOH (1154g, 20.56 moles), tetrabutylammonium hydrogen sulfate (140g, 0.411 mole), and deionized water (5.6L). The mixture was cooled to a temperature of approximately 20°C to 25°C. Methylene chloride (5.6L) was added to the flask, then 2-isopropylphenol (590g, 4.33 moles). The mixture was stirred for 0.5h before iodomethane (741g, 5.22 moles) was added. The stirring was continued at room temperature until the reaction was observed to be complete by HPLC, typically after approximately 5h. Stirring was discontinued and the reaction mixture was allowed to settle into aqueous and organic layers, which were then separated. Triethylamine (185 mL, 1.33 moles) was added to the organic phase with stirring for at least 15 min. Solvent was then removed by vacuum distillation to afford a white solid. Cyclohexane (4l) was added to the white solid, which was filtered and washed with further cyclohexane to remove the reaction product as a solvate. The cyclohexane solution was washed with aqueous HCl (2N), then half-saturated aqueous NaCl, and finally saturated aqueous NaCl. The cyclohexane was removed by vacuum distillation to afford 1-isopropyl-2-methoxybenzene as a light yellow oil of medium viscosity (612g, 94% yield).

EXAMPLE 2**Preparation of 3-isopropyl-4-methoxybenzaldehyde**

A 5L three-necked round bottom flask was equipped with mechanical agitation and provided with an inert gas blanket. The flask was charged with 1-isopropyl-2-

methoxybenzene (859g, 5.85 moles) and dimethylformamide (1584mL, 20.46 moles), and the mixture was heated to 80°C. Phosphorus oxychloride (2690g, 17.54 moles) was added to the reaction mixture over approximately 3h while maintaining the temperature of the reaction mixture between 80°C and 90°C. The addition of POCl₃ caused an immediate color change with continuous darkening over the course of the reaction. The reaction mixture was maintained at 80°C for approximately 16h, then checked by HPLC for presence of unreacted starting materials. The reaction mixture was quenched by slow addition over approximately 2h to icewater (8 kg total, 5 kg ice); alternatively the reaction mixture was added over approximately 3h to deionized water (10L) in a jacketed 30L reactor whose cooling bath was maintained between 0° and 5°C. Ethyl acetate (12L) was added to the quenched reaction with stirring. After settling, the layers were separated, and the aqueous layer was washed with additional ethyl acetate (4L). The combined organic layers were washed with saturated aqueous sodium bicarbonate solution (4L) and saturated aqueous sodium chloride (4L). The ethyl acetate was removed by vacuum distillation to yield 3-isopropyl-4-methoxybenzaldehyde (881g, 86.5%).

EXAMPLE 3

Preparation of hydroxy-(3-isopropyl-4-methoxy-phenyl)methanesulfonic acid, sodium salt

A 22L three-necked round bottom flask equipped with mechanical agitation was charged with 3-isopropyl-4-methoxybenzaldehyde (880g, 4.94 moles), THF (4650mL), and cyclohexane (3740mL) at room temperature. To this mixture was added, with stirring, an aqueous solution of

sodium bisulfite (1310g, 12.56 moles, in 4360mL of deionized water). The reaction mixture was stirred overnight at room temperature. The reaction product was removed by filtration, washed liberally with
5 cyclohexane:THF (3:1) and dried overnight by evaporation under air flow at room temperature to yield hydroxy-(3-isopropyl-4-methoxyphenyl)methanesulfonic acid as the sodium salt (1302g, 93.4%).

10

EXAMPLE 4**Preparation of 3-isopropyl-4-methoxyphenol**

A 22L three-necked round bottom flask equipped with mechanical agitation was charged with hydroxy-(3-isopropyl-4-methoxyphenyl)methanesulfonic acid, sodium
15 salt (1300g, 4.61 moles), *p*-toluenesulfonic acid monohydrate (908g, 4.77 moles), and methanol (10.4L), and the mixture was cooled to 0° to 5°C. Hydrogen peroxide (30% aqueous, 1.625L, 16.12 moles) was added slowly to the reaction mixture over about 2h. The reaction mixture
20 was stirred overnight and allowed to return to room temperature. The completeness of the reaction was assayed by HPLC; if the reaction was incomplete, further hydrogen peroxide was added to the reaction mixture. The reaction mixture was cooled once more to 0° to 5°C and
25 then quenched by addition of an aqueous slurry of sodium dithionite (1860g, 10.68 moles, stirred 1h at room temperature in 6.5 L deionized water). After stirring at least 0.5h, the reaction mixture was filtered, and the white solids were washed with ethyl acetate (8L). The
30 ethyl acetate and the aqueous filtrate were combined, agitated, and allowed to separate. Cyclohexane was added if necessary to facilitate separation. The organic phase was washed with sodium bicarbonate (4L, 10% aqueous

solution), then with saturated aqueous sodium chloride solution (4L). The solvent was removed by vacuum distillation to afford 3-isopropyl-4-methoxyphenol (510g, 66.7%).

5

EXAMPLE 5

Preparation of Trifluoromethanesulfonic acid 2,6-dibromo-4-nitrophenyl ester

10 A 12L jacketed vessel equipped with a mechanical stirrer and nitrogen inlet adapter was charged with dichloromethane (3.2L) and 2,5-dibromo-4-nitrophenol (800g, 2.69 moles). The reaction mixture was cooled to 0°C and sparged with nitrogen gas. Pyridine (0.436L, 3.23 moles) was introduced via addition funnel while the
15 temperature of the reaction mixture was maintained below 8°C. The reaction mixture was stirred at 0°C for 5 min, then a solution of triflic anhydride (0.544L, 3.23 moles) in dichloromethane (0.400L) was added via addition funnel while maintaining the reaction temperature below 10°C.
20 The reaction mixture was warmed to room temperature, stirred 30 min, and monitored periodically by HPLC. After the reaction had run to completion, aqueous hydrochloric acid (1N, 1.6L) was added to the reaction mixture dropwise with stirring. The reaction temperature
25 was maintained below 35°C during the addition of the acid. The reaction mixture was allowed to settle into layers, and the organic phase was separated and washed first with saturated aqueous sodium bicarbonate (0.8L), then with saturated aqueous sodium chloride (0.8L). The
30 organic phase was concentrated by rotary evaporation under vacuum at 40°C, then dried overnight under vacuum to yield trifluoromethane sulfonic acid 2,6-dibromo-4-nitrophenyl ester (1140g, 99.1%).

EXAMPLE 6**Preparation of 1,3-dibromo-2-iodo-5-nitrobenzene**

Dimethylformamide (4L) and trifluoromethane sulfonic
5 acid 2,6-dibromo-4-nitrophenyl ester (1135g, 2.65 moles)
were charged into a 22L jacketed vessel equipped with a
mechanical stirrer, a condenser, and a nitrogen purge.
Sodium iodide (1595g, 10.64 moles) was added to the
reaction mixture in portions while maintaining the
10 reaction temperature below 25°C. The reaction mixture
was then heated to 100°C and monitored for completion by
HPLC. Upon completion, the reaction mixture was cooled
to 0°C. Water (4L) was added while maintaining the
temperature below 14°C. The reaction mixture was stirred
15 for 30min, then filtered and washed with water (1.5L).
The resultant solid product was dried overnight by
evaporation under air flow at room temperature to yield
1,3-dibromo-2-iodo-5-nitrobenzene (892.7g, 82.9%). The
product may optionally be purified further by
20 recrystallization from ethanol.

EXAMPLE 7**Preparation of 4-(2,6-dibromo-4-nitrophenoxy)-
2-isopropylanisole**

25 A 22L jacketed vessel equipped with mechanical stirring
and a nitrogen inlet adapter was charged with
dimethylformamide (4L) and 3-isopropyl-4-methoxyphenol
(462.8g, 2.59 moles). The solution was cooled to 0°C,
and KHMDS (5.50L of 0.5M toluene solution, 2.25 moles)
30 was added over 1.5h while maintaining the temperature of
the mixture below 2°C. The reaction mixture was stirred
for 30 min at 2°C, then 1,3-dibromo-2-iodo-5-nitro-
benzene (798.6g, 1.96 moles) was added in portions, still

maintaining temperature of the mixture below 2°C. After the addition was complete, the reaction mixture was allowed to stir at room temperature for 2h or until the reaction was complete as assayed by HPLC. The reaction mixture was cooled once more to 2°C and ethyl acetate (4.5L) was added, followed by slow addition of saturated aqueous sodium bicarbonate (6L) while maintaining the temperature of the reaction mixture below 28°C. The organic layer was washed with saturated aqueous sodium chloride solution, then concentrated by rotary evaporation. The black, oily residue was taken up in acetone (3.87L). Water (0.77L) was added to the solution over 30 min with heating to 42°C. The solution was seeded with crystals of the product, cooled to 0°C, and stirred for 15 min. The resulting crystals of 4-(2,6-dibromo-4-nitrophenoxy)-2-isopropylanisole (665g, 76.1%) were recovered by filtration and dried for 48h by evaporation under air flow at room temperature.

20

EXAMPLE 7A

Preparation of 4-(2,6-dibromo-4-nitrophenoxy)- 2-isopropylanisole

3-Isopropyl-4-methoxyphenol (197.2g, 1.19 moles) was dissolved in dimethylformamide (4L). The mixture was stirred while the reaction vessel was cooled in an icewater bath. Sodium hydroxide (1.278L of 1.0M aqueous solution, 1.278 moles) was added while maintaining the temperature of the reaction mixture below 35°C. 1,3-Dibromo-2-iodo-5-nitrobenzene (400.0g, 0.983 moles) dissolved in dimethylformamide (portion of 4L) was added to the reaction mixture over approximately 45 min. The reaction mixture was stirred at 20° to 25°C for 24h. Deionized water (2L) was added over 2h and the reaction

mixture was cooled to 5°C. The resulting crystals of 4-(2,6-dibromo-4-nitrophenoxy)-2-isopropylanisole (405.4g, 92.6%) were recovered by filtration and dried for 48h by evaporation under air flow at room temperature.

5

EXAMPLE 8

Preparation of 4-(2,6-dibromo-4-nitrophenoxy)- 2-isopropylphenol

10 A 12L three-necked round bottom flask equipped with a mechanical stirrer, an addition funnel, and a temperature probe was charged with 4-(2,6-dibromo-4-nitrophenoxy)-2-isopropylanisole (752.1g, 1.69 moles), which was dissolved by stirring in dichloromethane (2L) at room temperature. The mixture was cooled to 8.7°C in
15 an ice-water bath. A solution of boron tribromide (2.050L, 1M in CH₂Cl₂, 2.05 moles) was added over 30 min. An increase of approximately 10°C in the temperature of the reaction mixture was observed. The ice-water bath was removed and the reaction mixture was allowed to stir
20 at room temperature until the reaction was complete, as assayed by HPLC. The reaction mixture was cooled once more in an ice-water bath, then quenched by slow addition of methanol (850mL), during which the temperature of the reaction mixture was maintained below 18°C. Saturated
25 aqueous sodium bicarbonate solution was added until the pH of the reaction mixture was 6 or 7. The aqueous phase was separated and washed with aliquots of ethyl acetate (2x1L). The organic layers were combined and the solvent removed under a vacuum at room temperature to yield crude
30 4-(2,6-dibromo-4-nitrophenoxy)-2-isopropylphenol (796g, 110%).

EXAMPLE 9

Preparation of 4-(4-amino-2,6-dibromophenoxy)-
2-isopropylphenol

4-(2,6-Dibromo-4-nitrophenoxy)-2-isopropylphenol
5 (795g, 1.84 moles) was divided into three roughly equal
portions. Each portion was dissolved in THF (2.6L) and
charged into a 12L three-necked round bottom flask
equipped with a mechanical stirrer, addition funnel, and
temperature probe. The solutions were heated to 33°C.
10 Sodium dithionite (482g, 2.4 moles) dissolved in
deionized water (2.6L) was added to each flask. The
reaction temperature was held at 50°C until each reaction
was complete as assayed by HPLC. Hydrochloric acid (6N
aqueous) was added dropwise to the heated reaction
15 mixtures until their pH was about 3, and heating was
continued for 45 min. The reaction mixtures were cooled
to room temperature and their pH was adjusted to 6.6 by
addition of saturated aqueous sodium bicarbonate
solution. Ethyl acetate (2.5L) was added to each flask
20 with stirring for 2 min. Each reaction mixture was
separated, and each organic layer was washed once with
half-saturated aqueous sodium chloride solution, then
once with saturated aqueous sodium chloride solution.
The solvents were removed by rotary evaporation and the
25 solids were dried overnight in a vacuum oven at 40°C
to yield crude 4-(4-amino-2,6-dibromophenoxy)-
2-isopropylphenol (657g, 96.9%). The crude 4-(4-amino-
2,6-dibromophenoxy)-2-isopropylphenol was purified by
recrystallization from isopropanol upon addition of 2.5
30 volumes of deionized water (90% recovery).

EXAMPLE 10

Preparation of *N*-[3,5-dibromo-4-(4-hydroxy-3-isopropylphenoxy)phenyl]malonamic acid

A slurry of 4-(4-amino-2,6-dibromophenoxy)-2-isopropylphenol (160g, 0.401 mole) in *t*-butyl methyl ether (TBME, 1.07L) was prepared in a 3L, three-necked round bottom flask equipped with a mechanical stirrer, a thermocouple probe and an addition funnel. The mixture was cooled to 2°C using a dry ice-acetone bath. A solution of sodium hydroxide (0.76M, 0.53L, 0.401 mole) was added over 20 min while keeping the internal temperature of the reaction mixture below 2°C. Ethyl malonyl chloride (78.2 mL, 0.614 mole) was added via addition funnel to the biphasic mixture over 10 min while still maintaining the internal temperature below 2°C. The cooling bath was removed and the mixture was stirred for 40 min at room temperature. Fifteen minutes after the addition of malonyl chloride was completed, the reaction temperature was 5°C and HPLC analysis of the reaction mixture failed to detect any remaining starting material. The reaction mixture was once more cooled to 2°C using a dry ice-acetone bath, and a solution of sodium hydroxide (10.0 M, 176 mL, 1.76 moles) was added over 10 min. The internal temperature of the reaction mixture was maintained below 2°C. After the addition was complete, the cooling bath was removed and the mixture was stirred at room temperature. Two hours after the addition of NaOH was complete, the reaction temperature was 19°C and none of the intermediate ethyl ester was detected by HPLC analysis. The organic layer was separated and extracted with water (150mL). Any remaining traces of organic solvent were removed from the combined aqueous layers by distillation under reduced

pressure. The aqueous solution was cooled in a dry ice-acetone bath to an internal temperature of 1°C and acidified with HCl (10% aqueous, 1.25L). The internal temperature of the mixture was maintained below 2°C throughout the addition. The resultant precipitated solid was collected by filtration and washed twice with water. The solid was dried to constant weight in a vacuum oven at 40°C to provide crude *N*-[3,5-dibromo-4-(4-hydroxy-3-isopropylphenoxy)phenyl]malonamic acid as a beige solid (219.5g, 113%). The crude product was recrystallized with 90% recovery from methanol upon addition of water and cooling to 0°C.

COMPARATIVE EXAMPLE

Acetic anhydride (9.9g, 97.5 mmol) was charged to a three-necked round bottom flask equipped with magnetic stirring, a nitrogen-inlet adapter, and an addition funnel. The acetic anhydride was cooled to a temperature of 20° to 25°C by immersing the flask in a dry ice-ethanol bath. Fuming nitric acid (6g, 94.9 mmol) was added dropwise to the acetic anhydride while maintaining the internal temperature of the mixture. Iodine (1g, 3.94 mmol) was added in one portion, followed by the addition of trifluoroacetic anhydride (5.4g, 47.3 mmol). The reaction mixture, which contained the unisolated intermediate iodonium salt (CF₃CO₂)₃I, was warmed to room temperature.

An aliquot of the reaction mixture (9mL) in a Reactive System Screening Tool (RSST) was placed inside a bomb. The bomb was pressurized with nitrogen (300 psig at 15°C) and heated at the rate of 1 C°/min. The 10 mL glass cell of the RSST did not break; however, some of

the reaction mixture erupted into the bomb, and the remainder of the aliquot changed color from red to dark purple-black.

5 Figure 1 is a graph of the resulting temperature vs. time measurements. It is apparent that an exothermic reaction began almost immediately with the commencement of heating. Similar results were obtained in differential scanning calorimetry (DSC) experiments. The
10 data in Figure 1, and the accompanying observations, illustrate that the thermal decomposition of the unisolated iodonium salt $(\text{CF}_3\text{CO}_2)_3\text{I}$ is so highly exothermic as to render $(\text{CF}_3\text{CO}_2)_3\text{I}$ hazardous as an intermediate in large scale preparations.

15 2-Isopropylphenol was reacted with methyl iodide in the presence of potassium carbonate to yield 1-isopropyl-2-methoxybenzene, which in turn was reacted with the solution of unisolated intermediate $(\text{CF}_3\text{CO}_2)_3\text{I}$ described
20 above to form the *bis*-[3-isopropyl-4-methoxyphenyl] iodonium salt in 40-70% yield. The *bis*-[3-isopropyl-4-methoxyphenyl] iodonium salt precipitated as a gummy solid, making purification impracticable and contributing to the unpredictability of the yield.

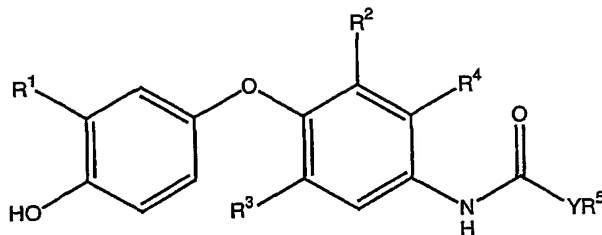
25 The *bis*-[3-isopropyl-4-methoxyphenyl] iodonium salt was reacted with 2,6-dibromo-4-nitrophenol in the presence of elemental copper and triethylamine, over the course of 4 to 5 days in the dark, to form 4-(4-nitro-
30 2,6-dibromophenoxy)-2-isopropylphenol in 50 to 80% yield. The extended time required for the ether formation, its unpredictable yield, and the unusual necessity to exclude

light are inefficiencies that also militate against commercial practice of this synthetic route.

5 The entire disclosures of the patents and other
publications cited above are incorporated herein by
reference. While certain preferred embodiments of the
present invention have been described and specifically
exemplified above, it is not intended that the invention
be limited to such embodiments. Various modifications
10 may be made to the invention without departing from the
scope and spirit thereof as set forth in the following
claims.

WHAT IS CLAIMED IS:

1. A process for synthesizing a compound of formula I



I

wherein

R¹ represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons;

R² and R³ are the same or different and represent hydrogen, halogen, an alkyl group of 1 to 4 carbons, or a cycloalkyl group of 3 to 6 carbons, at least one of R² and R³ being other than hydrogen;

R⁴ represents hydrogen or a lower alkyl group;

R⁵ represents a carboxylic acid or an alkyl ester thereof;

and

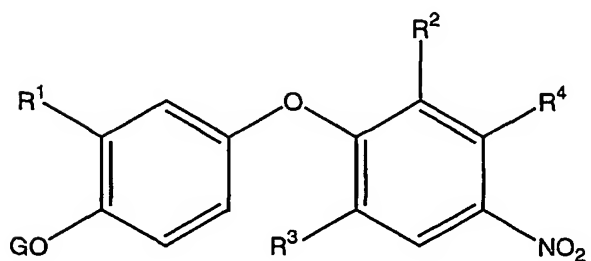
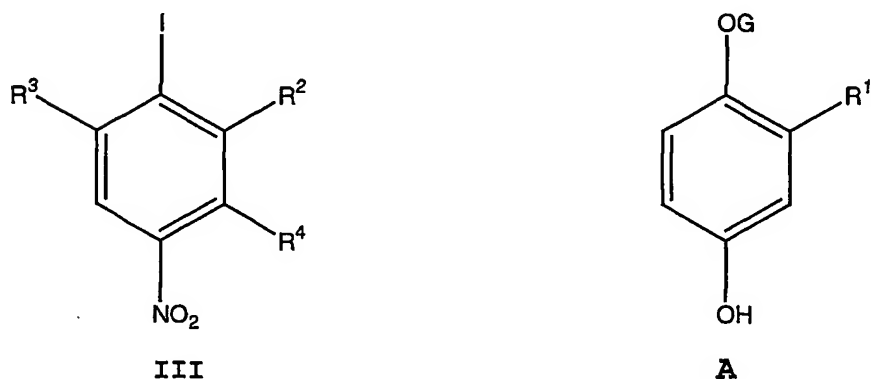
Y represents $-(CH_2)_n-$ where n is an integer from 1 to 5,

or a *cis*- or *trans*- ethylene group $-CH=CH-$;

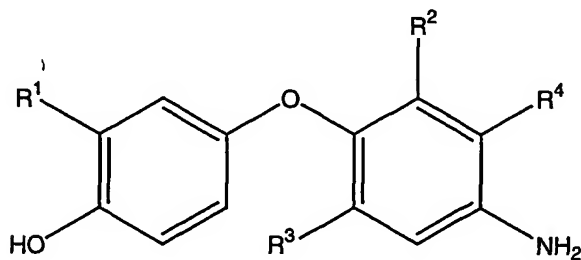
and all stereoisomers thereof;

said process comprising the steps of:

- a) reacting a compound of formula III with a phenol of formula A, wherein R¹, R², R³, and R⁴ are as defined above and G is a protecting group, in the presence of a base to produce a compound of formula IV;



- IV**
- b) deprotecting and reducing the compound of formula IV to form a compound of formula VI;



- c) reacting the compound of formula VI with a compound of the formula $\text{XC}(\text{O})\text{YR}^5$, wherein X

- represents OH or a halogen and R⁵ is a carboxylic acid alkyl ester, in the presence of a base, to produce a compound of formula I wherein R⁵ is a carboxylic acid alkyl ester; and
- 5 d) optionally de-esterifying the compound of formula I wherein R⁵ is a carboxylic acid alkyl ester to produce a compound of formula I wherein R⁵ is a carboxylic acid.
- 10 2. The process of claim 1, wherein, in step a, the base is selected from the group consisting of potassium hexamethyldisilazide (KHMDs), alkyl lithiums, lithium diisopropylamine, Grignard reagents, alkali metal hydrides, alkali metal carbonates, alkali
- 15 metal hydroxides, and alkali metal alkoxides.
3. The process of claim 2, wherein, in step a, the base is selected from the group consisting of potassium hexamethyldisilazide (KHMDs), potassium *t*-butoxide,
- 20 potassium *t*-pentoxide, LiHMDS, or NaHMDS
4. The process of claim 3, wherein, in step a, the base is potassium hexamethyldisilazide.
- 25 5. The process of claim 1, wherein, in step b, the deprotecting agent is selected from the group consisting of trialkylsilyl halides, mercaptans, boron halides, magnesium halides, aluminum halides, hydrogen over palladium catalyst, or iron halides.
- 30 6. The process of claim 5, wherein, in step b, the deprotecting agent is a trialkylsilyl halide or BBr₃.

7. The process of claim 6, wherein, in step b, the compound of formula IV is deprotected by reaction with BBr_3 .
- 5 8. The process of claim 1, wherein, in step b, the reducing agent is selected from the group consisting of elemental metals; hydrogen catalyzed by nickel, palladium or platinum; alkali metal borohydrides; magnesium-mediated reductions; iron carbonyls; iron
10 sulfate mediated reductions; iron and HCl; sodium hydrogen sulfite; sodium dithionite; tin and tin halide mediated reductions; and alkali metal sulfides and polysulfides.
- 15 9. The process of claim 8, wherein, in step b, the reducing agent is selected from the group consisting of sodium hydrogen sulfite; sodium dithionite; and iron and HCl.
- 20 10. The process of claim 9, wherein, in step b, the reducing agent is sodium dithionite.
11. The process of claim 1, wherein, in step c, the base
25 is selected from the group consisting of alkali metal hydroxides and alkoxides; alkaline earth metal hydroxides and alkoxides; alkyl amines; and substituted alkyl amines.
12. The process of claim 11, wherein, in step c, the
30 base is selected from the group consisting of alkali metal hydroxides and alkaline earth metal hydroxides.

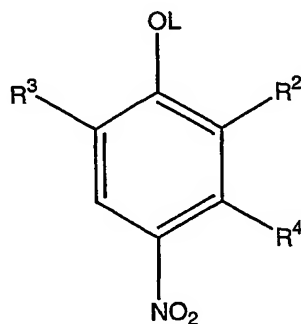
13. The process of claim 12, wherein, in step c, the base is sodium hydroxide.

14. The process of claim 1, wherein, in step d, the de-
esterifying agent is selected from the group
consisting of enzymes; hydrogen over catalytic
palladium; lithium and ammonia; zinc; alkali metal
hydroxides and alkoxides; alkaline earth metal
hydroxides and alkoxides; sodium sulfide; KSCN;
thioethers in combination with Lewis acids; anions
of alkyl thiols or selenols; trialkyl silyl halides;
zinc dichloride; mercury (II) acetate; and strong
protic acids.

15. The process of claim 14, wherein, in step d, the de-
esterifying agent is an alkali metal hydroxide.

16. The process of claim 15, wherein, in step d, the de-
esterifying agent is sodium hydroxide.

17. The process of claim 1, further comprising producing
the starting material of formula III by the step of
reacting a compound of formula II



II

wherein L is a leaving group;

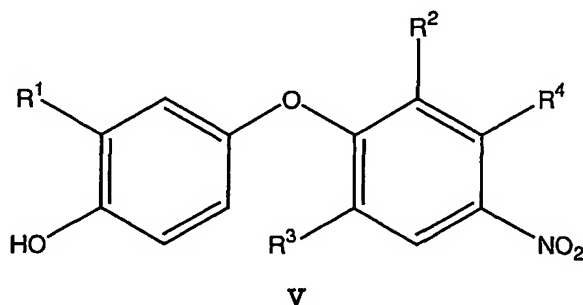
R^2 and R^3 are the same or different and represent hydrogen, halogen, an alkyl group of 1 to 4 carbons, or a cycloalkyl group of 3 to 6 carbons, at least one of R^2 and R^3 being other than hydrogen; and R^4 represents hydrogen or a lower alkyl group; with an iodinating agent.

18. The process of claim 17, wherein the iodinating agent is selected from the group consisting of alkali metal iodides and tetraalkylammonium iodides.

19. The process of claim 18, wherein the iodinating agent is selected from the group consisting of sodium iodide and potassium iodide.

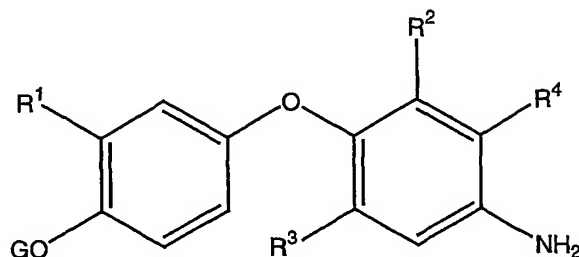
20. The process of claim 19, wherein the iodinating agent is sodium iodide.

21. The process of claim 1 wherein the compound of formula IV is first deprotected to form a compound of formula V



and then reduced to form the compound of formula VI.

22. The process of claim 1 wherein the compound of formula IV is first reduced to form a compound of formula VA



5

VA

and then deprotected to form the compound of formula VI.

23. The process of claim 1 for the preparation of a formula I compound wherein
 Y represents $-(CH_2)_n-$ where n is 1 or 2;
 R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons;
 R^2 and R^3 independently represent bromo, chloro, or methyl;
 R^4 represents hydrogen or methyl; and
 R^5 represents carboxyl.

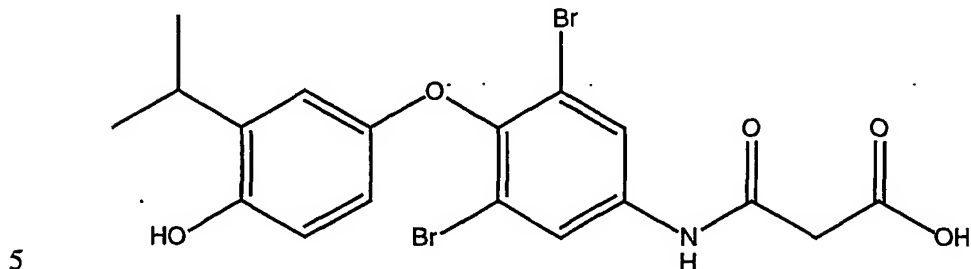
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24. The process of claim 1 for the preparation of a formula I compound wherein
 Y represents $-(CH_2)_n-$ where n is 1;
 R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons;
 R^2 and R^3 independently represent bromo or chloro;
 R^4 represents hydrogen or methyl; and

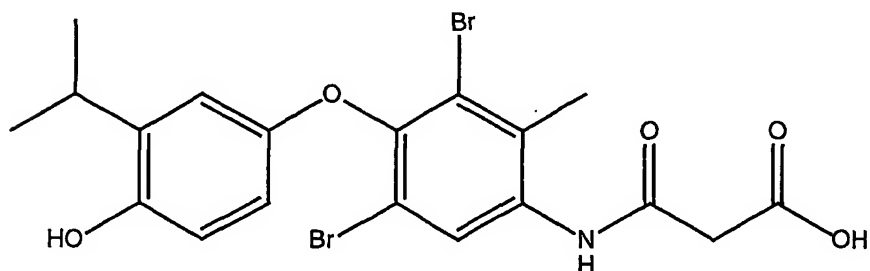
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R⁵ represents carboxyl.

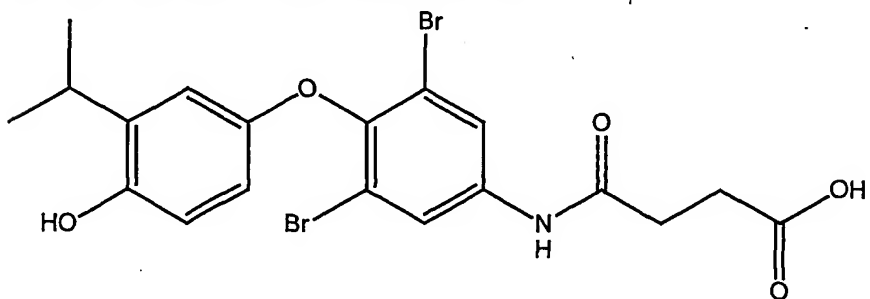
25. The process of claim 1 wherein the compound of formula I is selected from the group consisting of



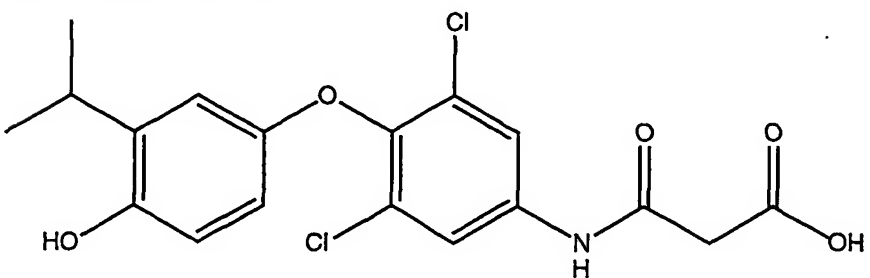
N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonamic acid;



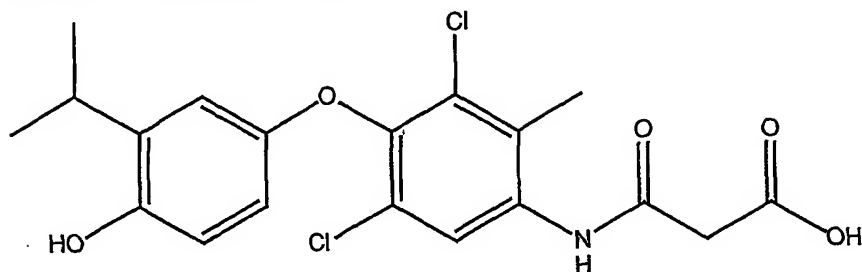
10 *N*-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-2-methyl-phenyl]-malonamic acid;



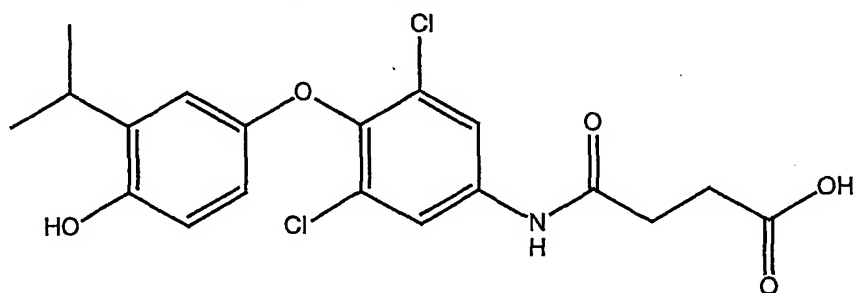
N-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-succinamic acid;



N-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonamic acid;

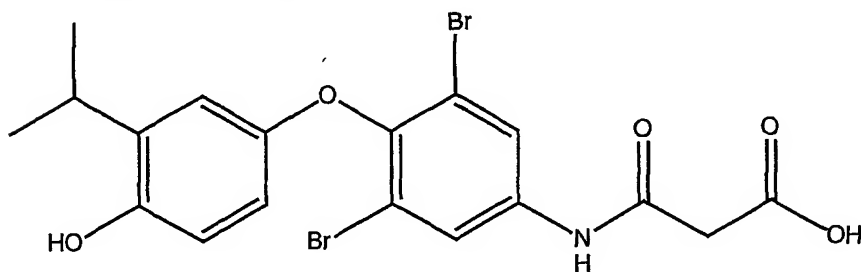


5 *N*-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-2-methyl-phenyl]-malonamic acid; and

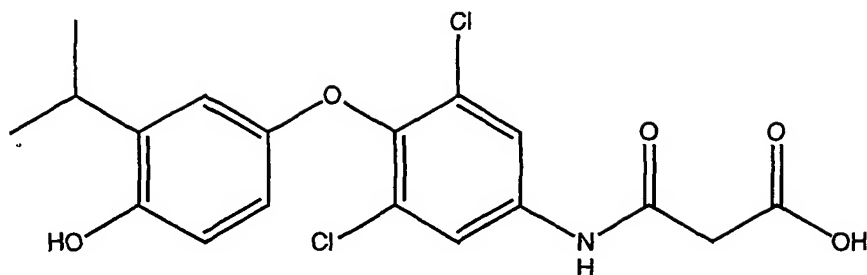


N-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-succinamic acid, and alkyl esters thereof.

10 26. The process of claim 1 wherein the compound of formula I is

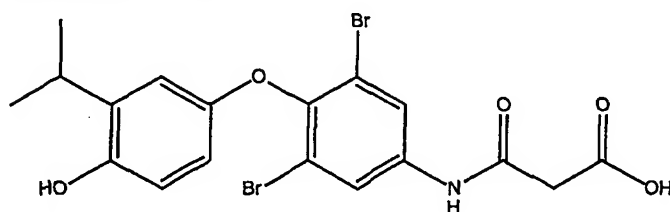


15 *N*-[3,5-dibromo-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonamic acid, or



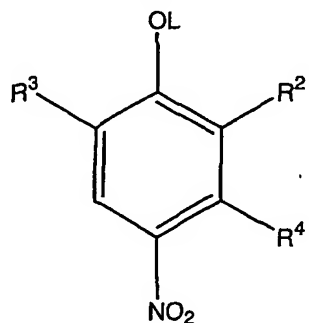
N-[3,5-dichloro-4-(4-hydroxy-3-isopropyl-phenoxy)-phenyl]-malonamic acid, and alkyl esters thereof.

- 5 27. The process of claim 1 wherein the compound of formula I is



and alkyl esters thereof.

- 10 28. A compound of formula IIA



IIA

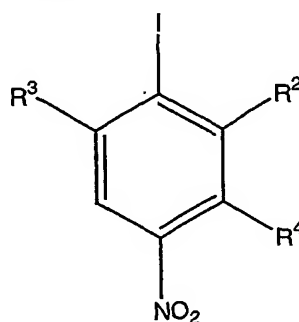
wherein

- 15 R^2 and R^3 independently represent bromo, chloro, or methyl;
 R^4 represents hydrogen or methyl; and
 L represents mesyl, tosyl, *p*-nitrobenzenesulfonyl, trihaloacetate, or triflate.

29. The compound according to claim 28, selected from the group consisting of trifluoro-methanesulfonic acid 2,6-dibromo-4-nitrophenyl ester, trifluoro-methanesulfonic acid 2,6-dibromo-3-methyl-4-nitrophenyl ester, trifluoro-methanesulfonic acid 2,6-dichloro-4-nitrophenyl ester, and trifluoro-methanesulfonic acid 2,6-dichloro-3-methyl-4-nitrophenyl ester.

30. The compound according to claim 28, trifluoro-methanesulfonic acid 2,6-dibromo-4-nitrophenyl ester.

31. A compound of formula IIIA



IIIA

wherein

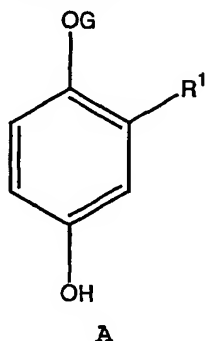
R² and R³ independently represent bromo, chloro, or methyl; and

R⁴ represents hydrogen or methyl.

32. The compound according to claim 31, selected from the group consisting of 1,3-dibromo-2-iodo-5-nitrobenzene, 1,3-dibromo-2-iodo-4-methyl-5-nitrobenzene, 1,3-dichloro-2-iodo-5-nitrobenzene, and 1,3-dichloro-2-iodo-4-methyl-5-nitrobenzene.

33. The compound according to claim 31, 1,3-dibromo-2-iodo-5-nitrobenzene.

5 34. A process for the synthesis of a phenol of formula A



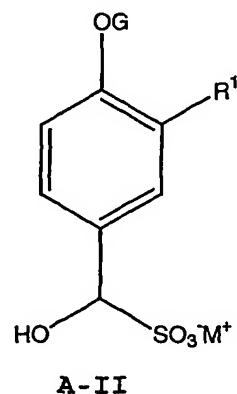
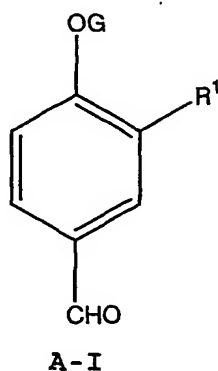
wherein

G is a protecting group; and

10 R^1 represents halogen, trifluoromethyl, an alkyl group of 1 to 6 carbons, or a cycloalkyl group of 3 to 7 carbons;

comprising the steps of:

15 a) reacting an aldehyde of formula A-I with MHSO_3 to form a sulfonate of formula A-II

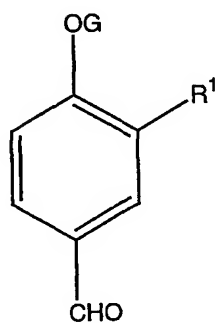


wherein M is a metal; and

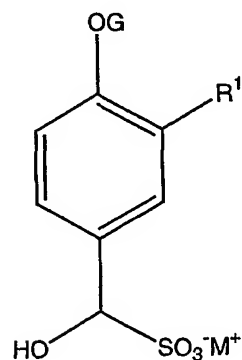
20 b) reacting the sulfonate of formula A-II with an oxidant, in the presence of a

proton source, to form the phenol of
formula A, above.

35. A compound selected from the group consisting of
5 compounds of formula A-I and formula A-II,



A-I



A-II

10

wherein

G represents a protecting group;

M represents a metal; and

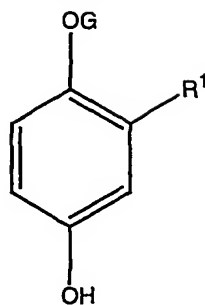
15 R^1 represents halogen, trifluoromethyl, an alkyl
group of 1 to 6 carbons, or a cycloalkyl group of 3
to 7 carbons.

36. The compound according to claim 35, 3-isopropyl-4-
methoxybenzaldehyde.

20

37. The compound according to claim 35, hydroxy-(3-
isopropyl-4-methoxyphenyl)methanesulfonic acid,
sodium salt.

38. A compound of formula A



A

5 wherein

 G represents a protecting group; and

 R¹ represents halogen, trifluoromethyl, an alkyl
 group of 1 to 6 carbons, or a cycloalkyl group of 3
 to 7 carbons;

10 with the proviso that, when G is a methyl group, R¹ is
 not a member of the group consisting of bromo, chloro,
 iodo, *t*-butyl, cyclohexyl, cyclopentyl, and isopropyl.

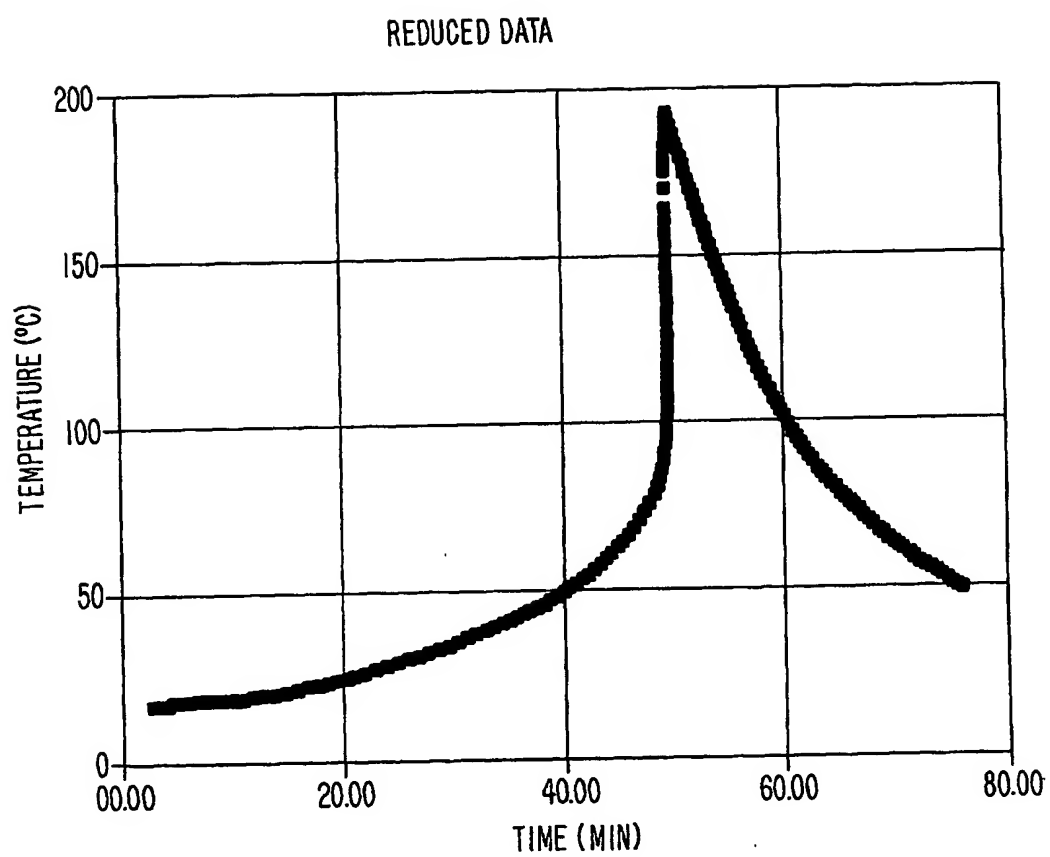


FIG. 1

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(74) Agents: BAXAM, Deanna et al.; Bristol-Myers Squibb Company, P.O. Box 4000, Princeton, NJ 08543-4000 (US).

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(71) Applicant (for all designated States except US): BRISTOL-MYERS SQUIBB COMPANY [US/US]; P. O. Box 4000, Route 206 and Provinceline Road, Princeton, NJ 08543-4000 (US).

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WO 03/039456 A3

(54) Title: PROCESS FOR THE PREPARATION OF ANILINE-DERIVED THYROID RECEPTOR LIGANDS

(57) Abstract: Provided are processes for the synthesis of aniline derivatives, specifically certain aniline derivatives which have activity as thyroid receptor ligands.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US02/34592

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07C 229/00

US CL : 560/42; 562/455

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 560/42; 562/455

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Please See Continuation Sheet**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y,P	US 2001/0051657 A1 (CHIANG et al) 13 December 2001 (13.12.2001), page 12, [0162]-pagepage 21, [0296], Scheme B, 1st two transformations.	1-3, 5-8, 11-16, 21-27
X	BAUM et al. Sulfonyl esters. 2. Carbon-Sulfur Bond Cleavage in Some Substitution Reactions of Nitrobenzenesulfonates. Canadian Journal of Chemistry. 1990, Vol 68, No. 8, pages 1450-1454, see page 1453, top, compound 7.	28, 29
X	BARBERO et al. 1-Aryl-3,3-dialkyltriazenes: A Convenient Synthesis from Dry Arenediazonium o-Benzenedisulfonimides - A High Yield Break Down to the Starting Dry Salts and Efficient Conversions to Aryl Iodides, Bromides and Chlorides. Synthesis. 2001, No.14, pages 2180-2190, see page 2188, Table 4, compounds 9k and 9l.	31
X	US 3,879,557 A (KAISER et al) 22 April 1975 (22.04.1975), column 4, lines 5-10.	35
X	Database CAPLUS on STN, AN 1968:95435. YAMANAKA et al. 'Chloromethylation of Phenol Derivatives'. Yakugaku Zasshi 1968, Vol 88, No.1, pages 116-117, Abstract, lines 7-8.	38
X	US 3,544,623 A (HANSEN et al) 01 December 1970 (01.12.1970), column 5, lines 24-61.	36

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"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

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INTERNATIONAL SEARCH REPORT

PCT/US02/34592

Continuation of B. FIELDS SEARCHED Item 3:

CAS ONLINE

search terms: structures, ?benzaldehyde, hydrogen sulfate, phenol, aryl iodide, aryl sulfonate

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